

# STIC Search Report

# STIC Database Tracking Number: 162359

TO: Jill M Gray

Location: REM 10A64

**Art Unit: 1774** 

September 2, 2005

Case Serial Number: 10/662350

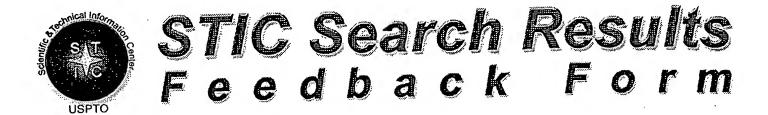
From: Les Henderson Location: EIC 1700 REM 4B28 / 4A30

Phone: 571-272-2538

Leslie.henderson@uspto.gov

Search Notes	• •	And the state of t		
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EIC17000

Questions about the scope or the results of the search? Contact the EIC searcher or contact:

Kathleen Fuller, EIC 1700 Team Leader 571/272-2505 REMSEN 4B28

Voluntary Results Feedback Form
<ul> <li>I am an examiner in Workgroup: Example: 1713</li> <li>Relevant prior art found, search results used as follows:</li> </ul>
102 rejection
103 rejection
Cited as being of interest.
Helped examiner better understand the invention.
Helped examiner better understand the state of the art in their technology.
Types of relevant prior art found:
☐ Foreign Patent(s)
Non-Patent Literature (journal articles, conference proceedings, new product announcements etc.)
> Relevant prior art not found:
Results verified the lack of relevant prior art (helped determine patentability).
Results were not useful in determining patentability or understanding the invention.
Comments:

Drop off or send completed forms to EIC1700 REMSEN 4B28



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Access DB# 16235

# SEARCH REQUEST FORM

Scientific and Technical Information Center

Requester's Full Name:Phone Mail Box and Bldg/Room Location	Number 30/2-1524	Examiner #: (06983 Date: 9/9/55 Serial Number: 0/662350 DAPER DISK E-MAIL
Please provide a detailed statement of the Include the elected species or structures,	****************** e search topic, and describe keywords, synonyms, acron s that may have a special me	as specifically as possible the subject matter to be searched.  ayms, and registry numbers, and combine with the concept or eaning. Give examples or relevant citations, authors, etc., if
Inventors (please provide full names):	Chers and the	s Use in the Preparation of Monwovers sanne; Mikelson, Torben; Mosler, Mikelson
Stengaard, Flemming	7/10/	<u> </u>
Earliest Priority Filing Date: 7	117/02	
appropriate serial number.	•••	parent, child, divisional, or issued patent numbers) along with the
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STAFF USE ONLY	**************************************	**************************************
Searcher: $\chi \pi$	NA Sequence (#)	STN \$ 1/5-4, 3 3
Searcher Phone #:	AA Sequence (#)	Dialog
Searcher Location:	Structure (#)	Questel/Orbit
Date Searcher Picked Up:	Bibliographic	Dr.Link
Date Completed: 9/1/05	Litigation	Lexis/Nexis
Searcher Prep & Review Time: 30	Fulltext	Sequence Systems
Clerical Prep Time:	Patent Family	WWW/Internet
Online Time: 230	Other	Other (specify)

PTO-1590 (8-01)

## AMENDMENTS TO THE CLAIMS

## List of Claims:

- (Previously Presented) A fibre comprising polyolefin polymer, said fibre having the features:
  - i) a fibre/fibre friction of no more than 600 g;
- ii) a spin finish consisting essentially of an aqueous emulsion of polysiloxanes, with at least 25% of the active content being polysiloxanes; and
  - iii) a fibre crystallinity of at least 50%.
- 2. (Original) A fibre according to claim 1 wherein the fibre/fibre friction is no more than 500 g.
- 3. (Original) A fibre according to claim 1 wherein the fibre/fibre friction is 200 to 600 g.
- 4. (Original) A fibre according to claim 1, wherein the spin finish consists essentially of an aqueous emulsion of polysiloxanes of at least 30% active content.
- 5. (Previously Presented) A fibre according to claim 4, wherein the spin finish is applied at a concentration of 2-15% wt/wt active content.

- 6. (Previously Presented) A fibre according to claim 4, wherein the spin finish level is 0.2 to 1% wt/wt with respect to the fibre.
- 7. (Original) A fibre according to claim 1, wherein the fibre crystallinity is at least 55% as measured by DSC or XRD.
- 8. (Original) A fibre according to claim 1, wherein the polyolefin polymer is a nucleated polymer.
- 9. (Previously Presented) A fibre according to claim 1, wherein the polyolefin polymer is a nucleated polymer, wherein the nucleating agent is selected from the group consisting of talc, metallic salts of aliphatic or aromatic carboxylic acids, branched polymers containing dendrittic branches and minerals selected from the group consisting of chalk, gypsum, clay kaolin, mica, and silicates and compounds that are based on D-sorbitol. 19
- 10. (Original) A fibre according to claim 9, wherein the nucleating agent is talc.

- 11. (Original) A fibre according to claim 9, wherein the polyolefin polymer is a nucleated polymer, nucleated with 5000 to 10000 ppm of talc. >(
- 12. (Original) A fibre according to claim 1, wherein the polyolefin is selected from the group consisting of isotactic or syndiotactic polypropylene homopolymers, homo and copolymers of monoolefins such as ethylene, propylene, alphaolefins, 4-methyl-1-pentene and blends thereof, linear polyethylenes, high density polyethylene, low density polyethylene, and linear low density polyethylene and blends of the same.
- 13. (Original) A fibre according to claim 9, wherein the polyolefin is selected from the group consisting of homopolymer polypropylene and homopolymer polyethylene.
- 14. (Original) A fibre according to claim 9, wherein the polyolefin is homopolymer polypropylene.
- 15. (Original) A fibre according to claim 1 with a bulk of at least about 30 cm<sup>3</sup>/g.

- 16. (Original) A fibre according to claim 1, wherein the draw ratio is about 1:2 to 1:8.
- 17. (Original) A fibre according to claim 1 having an ST dtex value of 2 to 20 dtex.
- 18. (Previously Presented) A fibre according to claim 1 having a resilience of at least about 40%.
- 19. (Previously Presented) A fibre according claim 1, wherein the polyolefin has a flexural modulus of at least 1500 MPa.

Claims 20-21 (Cancelled).

- 22. (Previously Presented) A fibre comprising polyolefin polymer according to claim 1, wherein the polyolefin polymer is a nucleated polymer, and said fibre has
  - i) a fibre/fibre friction of no more than 600 g;
- ii) a spin finish consisting essentially of an emulsion of polysiloxanes;

- iii) a draw ratio of at least 1:1.5 with a final fibre fineness of 2 to 10 dtex;
  - iv) a fibre crystallinity of at least 50%.
- 23. (Previously Presented) A non-woven material prepared from a polyolefin-based staple fibre as defined in any one of claims 1-19 and 22.
- 24. (Previously Presented) A non-woven material comprising polyolefin-based staple fibre, wherein the non-woven material has a bulk of at least 30 cm<sup>3</sup>/g and a resilience of at least 50%.
- 25. (Previously Presented) A non-woven material according to claim 24, wherein the non-woven material has a resilience of at least 55%.
- 26. (Previously Presented) A non-woven material according to any one of claims 24 to 25, wherein the nonwoven material has bulk of at least 35%.
- 27. (Currently Amended) A method of prepading a polyolefin-based fibre, said method characterised in the use of a nucleated polymer, a draw ratio of at least 1:1.5

with a final fibre dtex of 2 to 10 dtex., and a spin finish consisting essentially of an emulsion of polysiloxanes, with at least 25% of the active content being polysiloxanes.

- 28. (Previously Presented) A method according to claim 27, wherein the polymer is selected from polyethylene and polypropylene.
- 29. (Original) A method according to claim 27, wherein the draw ratio is 1:2 to 1:8.

Claim 30 (Cancelled),

- 31. (Currently Amended) A method according to claim 30 27, wherein the spin finish is applied at a concentration of 2-15% wt/wt active content.
- 32. (Currently Amended) A method according to claim 30 27, wherein the spin finish level is 0.2 to 1% wt/wt with respect to the fibre.
- 33. (Previously Presented) A method of preparing a non-woven material comprising the use of a fibre as defined

in any one claims 1 to 19 and 22, or the use of a fibre prepared according to the method according to any one of claims 27 to 32, comprising the steps of

- (a) forming a fibrous bond comprising said fibres, and
  - (b) bonding the fibrous web.
- 34. (Original) A method according to claim 33, wherein the fibres are oven-bonded at a temperature of 130 to 150 °C.
- 35. (Previously Presented) A fibre according to claim 1, wherein the fibre crystallinity of at least 50 % is achieved by:
  - iv) a draw ratio of at least 1:1.5; or
  - v) the polyolefin polymer being a nucleated polymer.
- 36. (Previously Presented) A fibre according to claim 1, wherein the spin finish is an external spin finish.

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(FILE 'HOME' ENTERED AT 14:27:18 ON.01 SEP 2005)

FILE 'HCAPLUS' ENTERED AT 14:27:34 ON 01 SEP 2005

E US20050079345/PN

L1 1 SEA ABB=ON PLU=ON US20050079345/PN SEL L1 RN

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FILE 'REGISTRY' ENTERED AT 14:28:07 ON 01 SEP 2005
             9 SEA ABB=ON PLU=ON (13397-24-5/BI OR 135861-56-2/BI OR
L2
               14807-96-6/BI OR 25085-53-4/BI OR 26063-22-9/BI OR
               50-70-4/BI OR 581798-19-8/BI OR 74-85-1/BI OR 9002-88-4/B
               I)
               D SCAN
               D L2 1-6 RN STR
               D L2 8-9 RN STR
               D L2 7 RN STR
L3
             1 SEA ABB=ON PLU=ON 13397-24-5/RN
               D SCAN
             1 SEA ABB=ON PLU=ON 135861-56-2/RN
L4
               D SCAN
L5
             1 SEA ABB=ON PLU=ON 14807-96-6/RN
               D SCAN
L6
             1 SEA ABB=ON PLU=ON 25085-53-4/RN
               D SCAN
L7
             1 SEA ABB=ON
                          PLU=ON 26063-22-9/RN
               D SCAN
             1 SEA ABB=ON PLU=ON 50-70-4/RN
L8
               D SCAN
L9
             1 SEA ABB=ON PLU=ON 581798-19-8/RN
               D SCAN
L10
             1 SEA ABB=ON PLU=ON 74-85-1/RN
               D SCAN
L11
             1 SEA ABB=ON
                          PLU=ON 9002-88-4/RN
               D SCAN
    FILE 'HCAPLUS' ENTERED AT 14:39:11 ON 01 SEP 2005
         27609 SEA ABB=ON PLU=ON L3
L12
L13
         45162 SEA ABB=ON PLU=ON GYPSUM?
L14
         46288 SEA ABB=ON
                          PLU=ON
                                  L12 OR L13
L15
           142 SEA ABB=ON
                          PLU=ON L4
L16
          2453 SEA ABB=ON
                          PLU=ON L15 OR D(A) SORBITOL? OR DSORBITOL?
L17
         29871 SEA ABB=ON
                          PLU=ON SORBITOL?
         27459 SEA ABB=ON · PLU=ON L17 NOT L16
L18
         29912 SEA ABB=ON
L19
                          PLU=ON L16 OR L17
L20
         24269 SEA ABB=ON
                           PLU=ON L5
L21
         43353 SEA ABB=ON
                          PLU=ON L20 OR TALC?
```

```
D 20-30 KWIC
L22
         19408 SEA ABB=ON PLU=ON L6
L23
         155787 SEA ABB=ON PLU=ON POLYPROPYLENE# OR (POLY O POLYM? OR
                HOMOPOLYM?) (A) PROPYLENE#
          17895 SEA ABB=ON PLU=ON ISOTACT? (2A) L23
L24
           1482 SEA ABB=ON PLU=ON SYNDIOTACT? (2A) L23
L25
L26
              8 SEA ABB=ON PLU=ON ADSTIF?
                D SCAN
          8857 SEA ABB=ON PLU=ON L23(2A)(COPOLYM? OR CO(W)POLYM?)
L27
          47662 SEA ABB=ON PLU=ON (COPOLYM? OR CO(W) POLYM?) (2A) PROPYLEN
L28
                E#
L29
          52761 SEA ABB=ON PLU=ON L27 OR L28
          17899 SEA ABB=ON PLU=ON L24 OR L26
L30
          1277 SEA ABB=ON PLU=ON L7
L31
L32
          1558 SEA ABB=ON PLU=ON L31 OR L25
L33
         185257 SEA ABB=ON PLU=ON (L23 OR L24 OR L25 OR L26 OR L27 OR
                L28 OR L29 OR L30 OR L31 OR L32)
          18952 SEA ABB=ON PLU=ON L8
L34
L35
           2325 SEA ABB=ON PLU=ON D(A) SORBITOL? OR DSORBITOL?
L36
              2 SEA ABB=ON PLU=ON L9
                D SCAN
           13 SEA ABB=ON PLU=ON L36 OR LUROL?
L37
          88596 SEA ABB=ON PLU=ON L10
L38
          31742 SEA ABB=ON PLU=ON ETHENE#
L39
          31026 SEA ABB=ON PLU=ON ALPHA(A)OLEFIN#
5872 SEA ABB=ON PLU=ON L39(L)L40
5748 SEA ABB=ON PLU=ON L39(3A)L40
L40
L41
L42
          5872 SEA ABB=ON PLU=ON L41 OR L42
L43
        173644 SEA ABB=ON PLU=ON L11
QUE ABB=ON PLU=ON POLYMER## OR HOMOPOLYMER## OR
L44
L45
                COPOLYMER## OR TERPOLYMER## OR RESIN? OR GUM?
L46
        2391109 SEA ABB=ON PLU=ON POLYETHYLENE# OR L45 OR ETHYLENE# (2A)
                L45
L47
             54 SEA ABB=ON PLU=ON ((FIBER?(A)FIBER?) OR (FIBRE#(A)FIBRE
                #))(2A)FRICTION?
              3 SEA ABB=ON PLU=ON G(2A)L47
L48
                D SCAN
                D 1-3 KWIC
L49
           3586 SEA ABB=ON PLU=ON (FIBER?(A)FIBER?) OR (FIBRE#(A)FIBRE#
                )
L50
             69 SEA ABB=ON PLU=ON L49(4A) FRICTION?
L51
             28 SEA ABB=ON PLU=ON L49(4A)G
                D 1-10 KWIC
L52
         741965 SEA ABB=ON PLU=ON E(W)G
             15 SEA ABB=ON PLU=ON L51 NOT L52
L53
                D L53 1-15 KWIC
L54
             13 SEA ABB=ON PLU=ON L51 NOT L53
                D 1-13 KWIC
```

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601 SEA ABB=ON PLU=ON SPIN? (2A) FINISH?
L55
L56
        1200984 SEA ABB=ON
                          PLU=ON FIBER? OR FIBR? OR FILAMENT? OR
               THREAD? OR STRAND? OR RIBBON? OR FILIFORM?
         117898 SEA ABB=ON PLU=ON POLYOLEFIN? OR OLEFIN? (2A) L45
         19871 SEA ABB=ON PLU=ON L56 AND L57
L58
L59
              2 SEA ABB=ON
                          PLU=ON L58 AND L50
               D SCAN
L60
             12 SEA ABB=ON PLU=ON L58 AND L55
               D SCAN
L61
             12 SEA ABB=ON PLU=ON L59 OR L60
L62
             1 SEA ABB=ON
                          PLU=ON L58 AND L53
               D SCAN
            12 SEA ABB=ON
                          PLU=ON L61 OR L62
L63
L64
         14949 SEA ABB=ON PLU=ON L56(2A)CRYST?
               D 1-9 KWIC
          2121 SEA ABB=ON PLU=ON L56(2A)CRYSTALLINIT?
L65
               D 5-9 KWIC
L66
             54 SEA ABB=ON
                          PLU=ON L65 AND L58
L67
             1 SEA ABB=ON
                          PLU=ON. L66 AND L55
               D SCAN
             1 SEA ABB=ON
                          PLU=ON L66 AND L50
L68
                           PLU=ON L66 AND L53
L69
             1 SEA ABB=ON
               D QUE L66
               D QUE L55
L70
        191279 SEA ABB=ON PLU=ON (SUSPEN? OR DISPERS? OR COLLOID? OR
               EMULS? OR MICROEMULS? OR SLURR?) (2A) (AQUEOUS? OR WATER?
               OR H2O)
             44 SEA ABB=ON
                          PLU=ON L55 AND L70
L71
L72
         155112 SEA ABB=ON PLU=ON ?SILOXANE?
             3 SEA ABB=ON PLU=ON L72 AND L71
L73
L74
          7065 SEA ABB=ON PLU=ON L72 AND L70
             3 SEA ABB=ON PLU=ON L74 AND L55
L75
               D QUE
L76
             O SEA ABB=ON PLU=ON L75 AND (PERCENT? OR PER(W)CENT? OR
               PCT?)
L77
             1 SEA ABB=ON PLU=ON L75 AND L58
               D OUE
               D QUE L70
L78
               QUE ABB=ON PLU=ON SUSPEN? OR DISPERS? OR COLLOID? OR
               EMULS? OR MICROEMULS? OR SLURR?
L79
           878 SEA ABB=ON PLU=ON L57 AND L78 AND L72
               D QUE
L80
             2 SEA ABB=ON
                          PLU=ON L79 AND L55
L81
             1 SEA ABB=ON
                          PLU=ON L79 AND L50
L82
             3 SEA ABB=ON PLU=ON L79 AND L49
            1 SEA ABB=ON
L83
                           PLU=ON
                                  L79 AND L65
            16 SEA ABB=ON PLU=ON L63 OR (L67 OR L68 OR L69) OR L73 OR
L84
               L75 OR L77 OR (L80 OR L81 OR L82 OR L83)
```

1	SEA ABB=ON SEA ABB=ON		ACTIV? (2A) CONTENT?
1			
		PLU=ON	(ACTIV?(2A)CONTENT?) AND L84
	D SCAN		
		PLU=ON	(ACTIV?(2A)CONTENT?) AND L58
			(ACTIV?(2A)CONTENT?) AND L55
-		120 011	(1101111 (211) (0111211111) 1212
18		PLH=ON	L88 OR L89
122			230 THE NOCHERIT.
. 37		PLII=ON	L91 AND L21
12			
29			
33/44			DSC:
72925			L95 OR DIFFER? (2A) SCAN? (2A) CALOR?
73033			195 OR DIFFER: (2A) SCAN: (2A) CABOR:
59793			XRD?
30793			ARD:
346409			(Y/W)DXY2\(/2X\DTEEDXCT2 OD 107 OD
340400		PHO-ON	(A(W)RAI:) (2A)DIFFRACI: OR H97 OR
0		DIJI_ON	194 AND /199 OD 196)
201		PLU=ON	102 MM (136 OK 136)
16		DI II_ON	L100 AND L58
10		PLU=ON	TIOO WID 128
4 5	-	DI II - ON	1101 OD 104
45		PLU=ON	L101 OR L94
2		DI II_ON	1102 AND 101
	•	PLO=ON	L102 AND L91
		DI II ON	1102 OD 1102
1		PTO=ON	L105 AND L104
4 =		DI 11 O1	110C OR 1104
			L108 AND METAL?
330896			DENDRIT? AND CHALK? OR KAOLIN? (2A) CLA
	· ·		
			(L110 OR L109) AND L105
			L110 AND L107
			L58 AND L110
			L58 AND L111
			L107 OR L112 OR L113 OR L115
			L116 AND (L50 OR L55 OR L65 OR L70)
17	SEA ABB=ON	PLU=ON	L116 AND L21
	3 18 1 18 122 37 12 29 53744 73835 58793 346408 0 581 16 45 2 45 7062 1 1 45 16414 6131 330896 1005 1 769 31 75 43	3 SEA ABB=ON 18 SEA ABB=ON 1 SEA ABB=ON D SCAN 18 SEA ABB=ON 122 SEA ABB=ON D QUE 37 SEA ABB=ON PCT? OR WT 29 SEA ABB=ON D L95 1-20 73835 SEA ABB=ON D L96 1-10 58793 SEA ABB=ON D L97 1-10 346408 SEA ABB=ON X (W) R (W) D 0 SEA ABB=ON D 1-9 KWIC 16 SEA ABB=ON D 1-9 KWIC 16 SEA ABB=ON D QUE 45 SEA ABB=ON D QUE 45 SEA ABB=ON D QUE 2 SEA ABB=ON D SCAN 45 SEA ABB=ON 1 SEA ABB=ON	3 SEA ABB=ON PLU=ON 18 SEA ABB=ON PLU=ON 1 SEA ABB=ON PLU=ON D SCAN 18 SEA ABB=ON PLU=ON 122 SEA ABB=ON PLU=ON D QUE 37 SEA ABB=ON PLU=ON PCT? OR WT OR WEIGH 29 SEA ABB=ON PLU=ON D L95 1-20 KWIC 73835 SEA ABB=ON PLU=ON D L96 1-10 KWIC 58793 SEA ABB=ON PLU=ON D L97 1-10 KWIC 346408 SEA ABB=ON PLU=ON X (W) R (W) D 0 SEA ABB=ON PLU=ON D L97 1-10 KWIC 346408 SEA ABB=ON PLU=ON D SCAN PLU=ON D SCAN PLU=ON D QUE 45 SEA ABB=ON PLU=ON D QUE 2 SEA ABB=ON PLU=ON D SCAN 45 SEA ABB=ON PLU=ON 1 SEA ABB=ON PLU=ON

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L119
             58 SEA ABB=ON PLU=ON L117 OR L118
L120
         265960 SEA ABB=ON PLU=ON PPM OR P(W)P(W)M OR PART? (2A)PER(2A)M
                ILLION?
L121
              O SEA ABB=ON PLU=ON L119 AND L120
         371683 SEA ABB=ON PLU=ON ?OLEFIN? OR L33
L122
                D SCAN L36
             49 SEA ABB=ON PLU=ON L119 AND (L122 OR (L38 OR L39 OR L40
L123
                OR L41 OR L42 OR L43 OR L44))
              1 SEA ABB=ON PLU=ON (L122 OR (L38 OR L39 OR L40 OR L41
L124
                OR L42 OR L43 OR L44)) AND L56 AND (L50 AND L55) AND
                (L65 OR L70)
               D SCAN
              2 SEA ABB=ON PLU=ON (L122 OR (L38 OR L39 OR L40 OR L41
L125
                OR L42 OR L43 OR L44)) AND L56 AND (L50 AND L55)
L126
             13 SEA ABB=ON PLU=ON (L122 OR (L38 OR L39 OR L40 OR L41
                OR L42 OR L43 OR L44)) AND L56 AND (L50 OR L55) AND (L65
                OR L70)
             56 SEA ABB=ON PLU=ON (L123 OR L124 OR L125 OR L126)
L127
L128
             5 SEA ABB=ON PLU=ON L127 AND BULK?
L129
             56 SEA ABB=ON PLU=ON L128 OR L127
L130
          22414 SEA ABB=ON PLU=ON CM(2A)G OR CENTIMET? (A) CUBE? (2A) (GRAM
                ? OR G) OR CENTIMET? (3A) (GRAM? OR G)
             O SEA ABB=ON PLU=ON L130 AND L129
L131
L132
             22 SEA ABB=ON PLU=ON L129 AND ((DRAW(2A)RATIO#) OR DTEX#
                OR RESILIENC? OR FLEX? OR MODUL? OR MPA OR M(W)P(W)A OR
               NON (W) WOVEN? OR NONWOVEN?)
L133
             56 SEA ABB=ON PLU=ON L129 OR L132
```

#### => => d l133 1-56 cbib abs hitstr hitind

L133 ANSWER 1 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN

- 2005:344315 Document No. 142:393463 Impact- and heat-resistant plastic
   moldings used in or on the ground. Matsuoka, Katsuhiro; Fujii,
   Kenji (Mikuni Plastics Co., Ltd., Japan; Daicel Polymer Ltd.). Jpn.
   Kokai Tokkyo Koho JP 2005105059 A2 20050421, 9 pp. (Japanese).
   CODEN: JKXXAF. APPLICATION: JP 2003-337786 20030929.
- AB The moldings comprise (A) 100 parts recycled PET and (B) 1-30 parts polyolefins containing (90-10):(10-90) (b-1) epoxy-containing olefin copolymers and (b-2) epoxy-free olefin copolymers. Thus, a composition containing recycled PET 100, Lotader AX 8840 (ethylene-glycidyl methacrylate copolymer) 9, and Lotryl 30BA02 (ethylene-Bu acrylate copolymer) 1 part was injection-molded to give a test piece showing Charpy impact strength (ISO 179) 6.5 kJ/m2, heat distortion temperature (ISO 75, under 1.8 MPa) 70°, and no sink mark.
- IT 14807-96-6, Talc, uses
  RL: MOA (Modifier or additive use); TEM (Technical or engineered

material use); USES (Uses)
 (filler; impact- and heat-resistant plastic moldings used in or
 on ground)
RN 14807-96-6 HCAPLUS
CN Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)

0 || HO-Si-OH

#### ●3/4 Mg

IC ICM C08L067-02 ICS C08K003-26; C08K003-34; C08K005-098; C08K007-04; G01F001-00; G01F015-18; C08L023-02

CC 38-3 (Plastics Fabrication and Uses) Section cross-reference(s): 60

ST plastic molding recycling PET **polyolefin** blend; ethylene glycidyl methacrylate butyl acrylate polymer blend; heat impact resistance plastic molding ground

IT Glass fibers, uses

RL: MOA (Modifier or additive use); TEM (Technical or engineered material use); USES (Uses)

(NEG-T 120; impact- and heat-resistant plastic moldings used in or on ground)

IT Mineral fibers

RL: MOA (Modifier or additive use); TEM (Technical or engineered material use); USES (Uses)

(filler; impact- and heat-resistant plastic moldings used in or on ground)

IT Mica-group minerals, uses

RL: MOA (Modifier or additive use); TEM (Technical or engineered material use); USES (Uses)

(fillers; impact- and heat-resistant plastic moldings used in or on ground)

IT Reinforced plastics

RL: TEM (Technical or engineered material use); USES (Uses) (glass fiber-reinforced; impact- and heat-resistant plastic moldings used in or on ground)

IT Crystal nucleating agents

Impact-resistant materials

Recycling of plastics and rubbers

(impact- and heat-resistant plastic moldings used in or on ground)

IT Polyesters, uses

Polyolefins

RL: POF (Polymer in formulation); TEM (Technical or engineered material use); USES (Uses)

(impact- and heat-resistant plastic moldings used in or on ground)

IT Alkali metal salts

Alkaline earth salts

RL: MOA (Modifier or additive use); TEM (Technical or engineered material use); USES (Uses)

(of organic carboxylic acids or carboxyl

-containing organic polymers, crystal nucleation

agents; impact- and heat-resistant plastic moldings used
in or on ground)

IT 822-16-2, AFCO-Chem NA

RL: MOA (Modifier or additive use); TEM (Technical or engineered material use); USES (Uses)

(crystal nucleation agent; impact- and

heat-resistant plastic moldings used in or on ground)

IT 471-34-1, Calcium carbonate, uses **14807-96-6**, **Talc** 

RL: MOA (Modifier or additive use); TEM (Technical or engineered material use); USES (Uses)

(filler; impact- and heat-resistant plastic moldings used in or on ground)

L133 ANSWER 2 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN

2005:340553 Document No. 142:393218 Regenerated PET compositions with improved moldability and heat and impact resistance, olefin resin compositions for modification of regenerated PET, and molded products of them. Fujii, Kenji (Daicel Polymer Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 2005105055 A2 20050421, 9 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 2003-337745 20030929.

One to 30 parts of olefin resin compns.

comprising epoxy group-containing olefin copolymers

and epoxy group-free olefin copolymers in the

weight ratio of 90:10 to 10:90 are added to 100 parts regenerated PET

to give the compns. Thus, recycled PET bottle flakes 100,

ethylene-glycidyl methacrylate copolymer (Lotader AX 8840) 9, and

ethylene-Bu acrylate copolymer (Lotryl 30BA02) 1 part were blended,

kneaded, pelletized, and injection molded to give test pieces

showing Charpy impact strength 6.5 kJ/m2, melt flow rate 5 g/10 min,

and heat distortion temperature 70°.

IT 14807-96-6, Talc, uses

RL: MOA (Modifier or additive use); USES (Uses) (reinforcement; regenerated PET-olefin resin

compns. with improved moldability and heat and impact resistance)

RN 14807-96-6 HCAPLUS

CN Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)

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#### ●3/4 Mq

IC ICM C08L067-02

ICS C08K003-26; C08K003-34; C08K005-098; C08K007-04; C08L023-08

CC 37-6 (Plastics Manufacture and Processing) Section cross-reference(s): 38

ST recycle PET glycidyl methacrylate ethylene polymer blend impact resistance; heat resistance regenerated PET polyolefin blend

IT Glass fibers, uses

RL: MOA (Modifier or additive use); USES (Uses)
(T 120, reinforcement; regenerated PET-olefin
resin compns. with improved moldability and heat and impact resistance)

IT Impact modifiers

Recycling of plastics and rubbers

(regenerated PET-olefin resin compns. with

improved moldability and heat and impact resistance)

IT Polyesters, properties

RL: POF (Polymer in formulation); PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(regenerated PET-olefin resin compns. with

improved moldability and heat and impact resistance)

IT Polymer blends

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(regenerated PET-olefin resin compns. with

improved moldability and heat and impact resistance)

IT Mica-group minerals, uses

Mineral fibers

RL: MOA (Modifier or additive use); USES (Uses)

(reinforcement; regenerated PET-olefin resin

compns. with improved moldability and heat and impact resistance)

IT 822-16-2, Sodium stearate

RL: MOA (Modifier or additive use); USES (Uses)

(crystal nucleating agent; regenerated PETolefin resin compns. with improved moldability

and heat and impact resistance)

IT 9010-86-0, Ethyl acrylate-ethylene copolymer 25103-74-6, Ethylene-methyl acrylate copolymer 25750-84-9, Lotryl 26061-90-5, Lotader AX 8840 29564-31-6, Ethylene-propyl acrylate copolymer 36604-80-5, Ethylene-glycidyl methacrylate-vinyl acetate copolymer 51541-08-3, Ethylene-glycidyl methacrylate-methyl acrylate copolymer 106677-58-1, Cevian V 680 RL: MOA (Modifier or additive use); USES (Uses) (regenerated PET-olefin resin compns. with improved moldability and heat and impact resistance) IT 25038-59-9, PET polymer, properties RL: POF (Polymer in formulation); PRP (Properties); TEM (Technical or engineered material use); USES (Uses) (regenerated PET-olefin resin compns. with improved moldability and heat and impact resistance) 471-34-1, Calcium carbonate, uses 14807-96-6, Talc IT , uses RL: MOA (Modifier or additive use); USES (Uses) (reinforcement; regenerated PET-olefin resin compns. with improved moldability and heat and impact resistance)

L133 ANSWER 3 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN

2004:680507 Document No. 141:175420 Polypropylene
fibers with low residual creep strain manufactured by melt
spinning polypropylene with degree of crystallinity
≥60%, drawing the fibers and heat-treating the
fibers and manufacture thereof. Fujii, Yasuyuki (Mitsubishi
Rayon Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 2004232117 A2
20040819, 8 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP
2003-20439 20030129.

AB The polypropylene (I) fibers (A1) have modulus 5-15 cN/dtex and show residual creep strain ≤20% on keeping the fibers for 5 h at stress 1 cN/dtex, or I fibers comprise above A1 fibers showing 2-point long-period image in the meridian direction as measured by the small angle x-ray diffractometry (SAXS) and exhibiting long period 10-20 nm. The I fibers are prepared by melt spinning I with melt flow rate (MFR) 10-50 g/10 min and degree of crystallinity ≥60%, drawing the fibers, and heat-treating the fibers for 5-60 s at 125-150° under dry hot air flow. I (TA-30) with MFR 30 g/10 min and degree of crystallinity 65% was melted at 200°, melt spun at nozzle temperature 200°, solidified in a cooling column, wound, drawn to draw ratio 2.7 at roll temperature 100°, and heat-treated at hot air temperature 140° for 5 s at fixed length to give 800-dtex/40filament fibers showing SAXS image and exhibiting modulus 7 cN/dtex and residual creep strain 9.8%.

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IC
     ICM D01F006-06
CC
     40-2 (Textiles and Fibers)
ST
     polypropylene fiber manuf creep strain redn;
     carpet polypropylene fiber creep strain redn
IT
     Polypropene fibers, uses
     RL: PEP (Physical, engineering or chemical process); PRP
     (Properties); PYP (Physical process); TEM (Technical or engineered
     material use); PROC (Process); USES (Uses)
        (polypropylene fibers with low residual creep
        strain manufactured by melt spinning polypropylene with
        specified degree of crystallinity, drawing the
        fibers and heat-treating the fibers and manufacture
        thereof)
ΙT
     Carpets
        (polypropylene fibers with low residual creep
        strain manufactured by melt spinning polypropylene with
        specified degree of crystallinity, drawing the
        fibers and heat-treating the fibers and manufacture
        thereof for)
IT
     25085-53-4, TA 30 (polyolefin)
     RL: PEP (Physical, engineering or chemical process); PRP
     (Properties); PYP (Physical process); TEM (Technical or engineered
     material use); PROC (Process); USES (Uses)
        (fiber; polypropylene fibers with
        low residual creep strain manufactured by melt spinning
        polypropylene with specified degree of
        crystallinity, drawing the fibers and
        heat-treating the fibers and manufacture thereof)
L133 ANSWER 4 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
2004:605461
             Document No. 141:158622 Improving the hydrophobicity of
     silicon-contg. vinyl polymers. Stark, Kurt; Herzig, Christian;
    Hoegl, Christian (Wacker Polymer Systems GmbH & Co. Kg, Germany).
    Ger. Offen. DE 10301976 A1 20040729, 19 pp.
                                                 (German). CODEN:
             APPLICATION: DE 2003-10301976 20030120.
AB
     Silicon-containing vinyl polymers with improved hydrophobicity for
    manufacture of coatings and plaster are manufactured by polymerization of
compns.
    containing (A) 60-99.99% silicon-containing monomer and ≥1 monomer
     selected from vinyl esters of C1-15 aliphatic carboxylic acids, C1-15
     alkyl (meth)acrylates, aromatic vinyl compds., olefins,
    dienes, and vinyl halides and (B) 0.01-40% branched polysiloxanes
    containing lipophilic siloxane structures and, optionally, linear or
    branched hydrophilic siloxane structures. A typical (B) was manufactured
    by heating a mixture containing 1,2,4-trivinylcyclohexane 108,
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 $\alpha, \omega$ -dihydrogen polymethylsiloxane ( active H

content 0.18%) 1840, and Karstedt catalyst 1.9 g 1 h at 80°, and heating the resulting intermediate (active H content 0.067%) 1 h at 100° with 3200 g monoallyl-terminated ethylene oxide-propylene oxide copolymer in the presence of a hexachloroplatinic acid solution

IC ICM C08G077-442

ICS C08L083-10; C09J183-10; C09D183-10

CC 42-10 (Coatings, Inks, and Related Products)

silicon contg vinyl polymer hydrophobicity enhancement branched siloxane additive; trivinylcyclohexane hydrogen polysiloxane allyl terminated polyoxyalkylene adduct manuf; olefin polymer silicon contg hydrophobicity enhancement branched siloxane additive; diene polymer silicon contg hydrophobicity enhancement branched siloxane additive; acrylic polymer silicon contg hydrophobicity enhancement branched siloxane additive; waterproof silicon contg vinyl polymer coating plaster

IT Nonwoven fabrics

(improving hydrophobicity of silicon-containing vinyl polymers by aqueous manufacture in presence of branched lipophilic silicon-containing polymers

as emulsifiers for binders for nonwoven fabrics)

IT Water-resistant materials

(improving hydrophobicity of silicon-containing vinyl polymers by aqueous manufacture in presence of branched lipophilic silicon-containing polymers

as emulsifiers for water-thinned plaster)

IT Fibers

RL: MSC (Miscellaneous)

(particulate materials; improving hydrophobicity of silicon-containing vinyl polymers by aqueous manufacture in presence of branched lipophilic silicon-containing polymers as emulsifiers for binders for particulate materials)

IT Coating materials

(water-resistant; improving hydrophobicity of silicon-containing vinyl polymers by aqueous manufacture in presence of branched lipophilic silicon-containing polymers as **emulsifiers** for water-thinned paints)

IT Adhesives

(water-thinned; improving hydrophobicity of silicon-containing vinyl polymers by aqueous manufacture in presence of branched lipophilic silicon-containing polymers as **emulsifiers** for water-thinned adhesives)

IT Paints

(water-thinned; improving hydrophobicity of silicon-containing vinyl polymers by aqueous manufacture in presence of branched lipophilic silicon-containing polymers as emulsifiers for water-thinned paints)

IT Plaster

(water-thinned; improving hydrophobicity of silicon-containing vinyl polymers by aqueous manufacture in presence of branched lipophilic

silicon-containing polymers as emulsifiers for
water-thinned plaster)

IT 2855-27-8DP, 1,2,4-Trivinylcyclohexane, reaction products with
hydrogen polysiloxanes and polyoxyalkylene monoallyl ether
9041-33-2DP, Ethylene oxide-propylene oxide
copolymer monoallyl ether, reaction products with
hydrogen-terminated methylsiloxanes and trivinylcyclohexane
RL: IMF (Industrial manufacture); MOA (Modifier or additive use);
TEM (Technical or engineered material use); PREP (Preparation); USES
(Uses)

(improving hydrophobicity of silicon-containing vinyl polymers by aqueous manufacture in presence of branched lipophilic silicon-containing polymers

as emulsifiers)

L133 ANSWER 5 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN 2004:515759 Document No. 141:55706 Process for making and process for converting polyolefin fibers. Smit, Leo; Simmelink, Joseph Arnold Paul Maria; Nielaba, Leonard Josef Arnold (DSM IP Assets B.V., Neth.). PCT Int. Appl. WO 2004053212 A1 20040624, 22 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, (English). CODEN: PIXXD2. APPLICATION: WO 2003-NL872 TG, TR. 20031209. PRIORITY: EP 2002-80143 20021210.

AB The invention relates to a process for making a multifilament polyethylene yarn via a gel-spinning process, wherein a spin finish is applied at least once in an amount of 0.1-10% based on the filament, to a filament that contains less than 50% of solvent; the spin finish comprising at least 95% of at least one volatile compound having a b.p. at 0.1 MPa pressure of from 30 to 250°; and the spin finish is subsequently removed by exposing the filament to a temperature of below the melting temperature of the filament. With this process a yarn is made that has a very low amount of residues on the surface of the fibers, without the need for a washing or extraction step, showing good mech. properties, and very suited for e.g. biomedical applications. The invention further relates to a process for converting polyolefin fibers into a semi-finished or end-use product. The invention also concerns a polyethylene yarn and a semi-finished or end-use product obtainable by said processes, as well as to the use of thereof in biomedical

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applications.
IT
     9002-88-4, Polyethylene
     RL: TEM (Technical or engineered material use); USES (Uses)
        (fibers; process for making polyolefin
        fibers with min. residual finish substance)
RN
     9002-88-4 HCAPLUS
     Ethene, homopolymer (9CI) (CA INDEX NAME)
CN
     CM
     CRN 74-85-1
     CMF C2 H4
H_2C = CH_2
     ICM D01F006-04
IC
CC
     40-2 (Textiles and Fibers)
ST
     gel spinning volatile compd finishing
     polyolefin yarn manuf; multi filament polyethylene
     yarn gel spinning volatile compd finishing
     Polyoxyalkylenes, uses
IT
     RL: REM (Removal or disposal); TEM (Technical or engineered material
     use); PROC (Process); USES (Uses)
        (derivs., finish, removal of; process for making
        polyolefin fibers with min. residual finish
        substance)
IT
     Polyolefin fibers
     RL: PRP (Properties); TEM (Technical or engineered material use);
     USES (Uses)
        (process for making polyolefin fibers with
        min. residual finish substance)
IT
     Alcohols, uses
     Ketones, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (volatile finish; process for making polyolefin
        fibers with min. residual finish substance)
IT
     9002-88-4, Polyethylene
     RL: TEM (Technical or engineered material use); USES (Uses)
        (fibers; process for making polyolefin
        fibers with min. residual finish substance)
IT
     25322-68-3D, Polyethylene oxide, derivs.
     RL: REM (Removal or disposal); TEM (Technical or engineered material
     use); PROC (Process); USES (Uses)
        (finish, removal of; process for making polyolefin
        fibers with min. residual finish substance)
IT
     118240-87-2, Isopropanol-water mixture 153523-39-8,
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1-Butanol-ethanol-water mixture RL: NUU (Other use, unclassified); USES (Uses) (volatile finish; process for making polyolefin fibers with min. residual finish substance)

L133 ANSWER 6 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN Document No. 142:199796 Crystallinity, orientation and 2004:488372 mechanical property development in melt spinning poly(ethylene-co-octene) filaments. Shan, Haifeng; White, James L. (Institute of Polymer Engineering, University of Akron, Akron, OH, 44325, USA). Annual Technical Conference - Society of Plastics Engineers, 62nd(Vol. 2), 2067-2071 (English) 2004. CODEN: ACPED4. ISSN: 0272-5223. Publisher: Society of Plastics Engineers. The structure development in melt spinning of poly(ethylene-co-AB octene) with different octene contents was studied. The filaments are characterized by WAXD, birefringence and DSC. Crystal structures of 4 copolymers are determined The Herman-Stein crystalline orientation factors are then calculated The uniaxial mech. properties are also measured.

CC 40-4 (Textiles and Fibers)

ITBirefringence Crystal structure Crystallinity Elongation at break Tensile strength

> (crystallinity and orientation and mech. property development in melt spinning poly(ethylene-co-octene) filaments)

IT Polyolefin fibers

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

(ethylene-octene; crystallinity and orientation and mech. property development in melt spinning poly(ethylene-cooctene) filaments)

IT Polymer chains

(orientation; crystallinity and orientation and mech. property development in melt spinning poly(ethylene-co-octene) filaments)

IT 70800-37-2, Ethylene-octene copolymer RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP (Physical process); PROC (Process) (fibers; crystallinity and orientation and mech. property development in melt spinning poly(ethylene-cooctene) filaments)

L133 ANSWER 7 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN Document No. 142:115811 Preliminary test of 2004:385453 electroconductive polyolefin fibers modified with polyaniline. Wlochowicz, A.; Fryczkowski, R. (Institute of

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Textile Engineering and Polymer Materials, University of
     Bielsko-Biata, Bielsko-Biala, 43-309, Pol.). Fibres & Textiles in
     Eastern Europe, 11(4), 36-38 (English) 2003. CODEN: FTEEEM.
     1230-3666. Publisher: Institute of Chemical Fibres.
     Initial research into methods for obtaining anti-static
AB
     fibers is presented. We used mixts. of
     polypropylene, serving as a polymer matrix, as well as
    polyaniline (PANI) as a conductive polymer and alkylbenzene sulfonic
     acid and a protonating agent. The concentration of PANI used in our
     research varied from 0 to 10%, and that of alkylbenzene sulfonic
     acid from 0 up to 30%. Polypropylene was added to
     complete the mixture up to 100% DSC and WAXS tests were
     carried out, as were elec. resistance measurements by the
     four-conductor connection technique. The results showed that in
     general, the melt mixts. were fiber-grade, and of breaking
     strength partly even 60 MPa. Electrocond. values of the
     order of 10-4 S/cm allowed the fibers obtained to be
     applied in fabrics with anti-static properties.
CC
     40-2 (Textiles and Fibers)
     Section cross-reference(s): 37, 76
    polypropylene fiber polyaniline alkylbenzene
ST
     sulfonic acid melting spun; fiber elec cond
     crystallinity
IT
     Polypropene fibers, uses
     RL: PEP (Physical, engineering or chemical process); POF (Polymer in
     formulation); PRP (Properties); PYP (Physical process); PROC
     (Process); USES (Uses)
        (Eltex P-HY 001P; electroconductive polyolefin
        fibers modified with polyaniline)
IT
    Breaking strength
    Conducting polymers
       Crystallinity
    Electric conductivity
    Melting
    Thermal stability
        (electroconductive polyolefin fibers modified
        with polyaniline)
IT
    Polyanilines
    RL: POF (Polymer in formulation); PRP (Properties); USES (Uses)
        (electroconductive polyolefin fibers modified
        with polyaniline)
IT
    Fibers
    RL: PEP (Physical, engineering or chemical process); PRP
     (Properties); PYP (Physical process); PROC (Process)
        (spinning; electroconductive polyolefin fibers
        modified with polyaniline)
IT
    Polymer morphology
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(surface; electroconductive polyolefin fibers

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modified with polyaniline)
IT
     25233-30-1, Polyaniline
     RL: POF (Polymer in formulation); PRP (Properties); USES (Uses)
        (electroconductive polyolefin fibers modified
        with polyaniline)
IT
     7440-57-5, Gold, uses
     RL: DEV (Device component use); USES (Uses)
        (electrode; electroconductive polyolefin fibers
        modified with polyaniline)
     25085-53-4, Isotactic polypropylene
IT
     RL: PEP (Physical, engineering or chemical process); POF (Polymer in
     formulation); PRP (Properties); PYP (Physical process); PROC
     (Process); USES (Uses)
        (fiber; electroconductive polyolefin
        fibers modified with polyaniline)
     7727-54-0, Ammonium persulfate
IT
     RL: CAT (Catalyst use); USES (Uses).
        (polymerization catalyst; electroconductive polyolefin
        fibers modified with polyaniline)
ΙT
     98-11-3D, Benzenesulfonic acid, alkyl derivs.
     RL: MOA (Modifier or additive use); USES (Uses)
        (protonating agent; electroconductive polyolefin
        fibers modified with polyaniline)
L133 ANSWER 8 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
             Document No. 140:288798 Polyolefin
2004:270153
     fibers and nonwoven fabrics with high bulk
     and resilience preparing from the same. Thomsen, Susanne
     Dahl; Mikkelsen, Torben Laigaard; Moller, Mikael; Stengaard,
     Flemming Faurby (Fibervisions A/S, Den.). PCT Int. Appl. WO
     2004027130 A1 20040401, 30 pp. DESIGNATED STATES: W: AE, AG, AL,
     AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ,
    DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID,
     IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA,
    MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU,
     SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
     VC, VN, YU, ZA, ZM, ZW; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY,
    DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT,
     SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO
     2003-DK600 20030917. PRIORITY: DK 2002-1368 20020917.
AB
     The nonwoven materials with bulk ≥30
     cm3/g and resilience ≥50% is prepared from
    polyolefin-based fibers ≥1 features
     selected from (i) fiber/fiber friction
     ≤600 g, (ii) spin finish
     comprising an aqueous emulsion of
    polysiloxane with active content
     ≥25%, (iii) draw ratio at least 1:1.5,
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and (iv) fiber crystallinity ≥50%. Thus,
     a nonwoven fabric obtained from fibers of
     Adstif HA 840R (nucleated polypropylene)
     treated with Silastol GF 602c (polysiloxane) in the
     drawing process was finished with Synthesin 7490 Fill, showing
     bulk 65 cm3/g and resiliency 74%.
IT
     74-85-1D, Ethene, polymers with
     \alpha -olefins
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); TEM (Technical or engineered material use); PROC
     (Process); USES (Uses)
        (LLDPE, fibers; polyolefin fibers
        and nonwoven fabrics with high bulk and
        resilience preparing from the same)
RN
     74-85-1 HCAPLUS
CN
     Ethene (9CI) (CA INDEX NAME)
H_2C = CH_2
IT
     13397-24-5, Gypsum, uses 14807-96-6,
     Talc, uses 135861-56-2, 1,3:2,4-Bis(3,4-
     dimethylbenzylidene) sorbitol
     RL: TEM (Technical or engineered material use); USES (Uses)
        (crystal nucleating agents;
        polyolefin fibers and nonwoven
        fabrics with high bulk and resilience preparing
        from the same)
RN
     13397-24-5 HCAPLUS
CN
     Gypsum (Ca(SO4).2H2O) (9CI) (CA INDEX NAME)
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Ca

●2 H<sub>2</sub>O

RN 14807-96-6 HCAPLUS CN Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)

### ●3/4 Mg

RN 135861-56-2 HCAPLUS
CN D-Glucitol, 1,3:2,4-bis-O-[(3,4-dimethylphenyl)methylene]- (9CI)
(CA INDEX NAME)

Absolute stereochemistry.

IT 9002-88-4, Polyethylene 26063-22-9,
 Polypropylene, syndiotactic
 RL: PEP (Physical, engineering or chemical process); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)
 (fibers; polyolefin fibers and nonwoven fabrics with high bulk and resilience preparing from the same)
RN 9002-88-4 HCAPLUS

CN Ethene, homopolymer (9CI) (CA INDEX NAME)

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CM
          1
     CRN 74-85-1
     CMF
         C2 H4
H_2C = CH_2
     26063-22-9 HCAPLUS
RN
     1-Propene, homopolymer, syndiotactic (9CI) (CA INDEX NAME)
     CM
     CRN 115-07-1
     CMF C3 H6
H_3C-CH=CH_2
IC
     ICM D01F006-04
         D01F006-30; D01F006-46; D04H001-00; D06M015-37; D06M015-643;
          C08F010-00
CC
     40-10 (Textiles and Fibers)
ST
     nonwoven fabric polyolefin fiber
     bulk resilience; polysiloxane
     spin finish polyolefin fiber
IT
     Polysiloxanes, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (Silastol GF 602c, finishing agent; polyolefin
        fibers and nonwoven fabrics with high
       bulk and resilience preparing from the same)
IT
     Silicone rubber, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (Synthesin 7490 Fill, finishing agent; polyolefin
        fibers and nonwoven fabrics with high
       bulk and resilience preparing from the same)
IT
     Fabric finishing
        (agents; polyolefin fibers and
       nonwoven fabrics with high bulk and
       resilience preparing from the same)
IT
     Chalk
      Dendritic polymers
     Kaolin, uses
      Mica-group minerals, uses
      Silicates, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
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(crystal nucleating agents;
        polyolefin fibers and nonwoven
        fabrics with high bulk and resilience preparing
        from the same)
ΙT
     Polyolefin fibers
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); TEM (Technical or engineered material use); PROC
     (Process); USES (Uses)
        (ethylene copolymers; polyolefin fibers and
        nonwoven fabrics with high bulk and
        resilience preparing from the same)
ΙT
     Polyolefin fibers
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); TEM (Technical or engineered material use); PROC
     (Process); USES (Uses)
        (ethylene; polyolefin fibers and
        nonwoven fabrics with high bulk and
        resilience preparing from the same)
IT
     Antistatic agents
     Crystal nucleating agents
       Nonwoven fabrics
        (polyolefin fibers and nonwoven
        fabrics with high bulk and resilience preparing
        from the same)
IT
    Polyolefin fibers
     Polypropene fibers, uses
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
    process); TEM (Technical or engineered material use); PROC
     (Process); USES (Uses)
        (polyolefin fibers and nonwoven
        fabrics with high bulk and resilience preparing
        from the same)
IT
    Carboxylic acids, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (salts, crystal nucleating agents;
        polyolefin fibers and nonwoven
        fabrics with high bulk and resilience preparing
        from the same)
IT
     25085-53-4, Adstif HA 840R
    RL: PEP (Physical, engineering or chemical process); PYP (Physical
    process); TEM (Technical or engineered material use); PROC
     (Process); USES (Uses)
        (Adstif HA 840R, nucleated, fiber;
       polyolefin fibers and nonwoven
        fabrics with high bulk and resilience preparing
        from the same)
IT
    74-85-1D, Ethene, polymers with
    \alpha -olefins
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RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); TEM (Technical or engineered material use); PROC
     (Process); USES (Uses)
        (LLDPE, fibers; polyolefin fibers
        and nonwoven fabrics with high bulk and
        resilience preparing from the same)
ΙT
     50-70-4D, D-Sorbitol, derivs. 13397-24-5
     , Gypsum, uses 14807-96-6, Talc, uses
     135861-56-2, 1,3:2,4-Bis(3,4-dimethylbenzylidene)
     sorbitol
     RL: TEM (Technical or engineered material use); USES (Uses)
        (crystal nucleating agents;
        polyolefin fibers and nonwoven
        fabrics with high bulk and resilience preparing
        from the same)
İT
     9002-88-4, Polyethylene 26063-22-9,
     Polypropylene, syndiotactic
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); TEM (Technical or engineered material use); PROC
     (Process); USES (Uses)
        (fibers; polyolefin fibers and
        nonwoven fabrics with high bulk and
        resilience preparing from the same)
IT
     581798-19-8, Lurol PP 920
     RL: NUU (Other use, unclassified); USES (Uses)
        (polyolefin fibers and nonwoven
        fabrics with high bulk and resilience preparing
        from the same)
L133 ANSWER 9 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
2003:998542
             Document No. 141:208456 Effects of on-line melt blending
     of polypropylene with polyamide 6 on the bulk
     and strength of the resulting BCF yarn. Tavanai, Hossein; Morshed,
    Mohammad; Hosseini, Seyed Majid (Textile Engineering Department,
     Isfahan University of Technology, Esfahan, 84156, Iran).
     Polymer Journal, 12(5), 421-430 (English) 2003. CODEN: IPJOFF.
     ISSN: 1026-1265. Publisher: Iran Polymer and Petrochemical
     Institute.
    Polypropylene filament yarns, when compared with
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AB Polypropylene filament yarns, when compared with polyethyleneterephthalate and polyamide yarns have lower stretchability after being false twist textured. To overcome this deficiency, other researchers have tried to lower the degree of crystallinity of this fiber by either tension annealing or blending it with polyethyleneterephthalate and polystyrene. These phys. or chemical modifications have not yet achieved a satisfactory level of enhancement of the texture-ability of polypropylene. Considering the importance of BCF polypropylene yarns, in this research, the effect of

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Temperature

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blending polypropylene with 12.5 and 25 % of polyamide 6
on the bulk of the final bulked continuous
filament (BCF) yarn has been investigated.
                                            The results show
that although both crimp contraction and crimp modulus
increase with the temperature and pressure of hot air during texturing,
but blending polypropylene with even 25% of polyamide 6
does not enhance the texture-ability of BCF yarns. Also, the reduction
in the crystallinity of the yarns does not lead to any improvement
in the texture-ability of the polypropylene BCF yarns.
40-4 (Textiles and Fibers)
Section cross-reference(s): 36
polypropylene polyamide blend yarn property crystallinity
texture ability
Contraction (mechanical)
  Modulus (stress-strain)
   (crimp; effects of online melt blending of polypropylene
   with polyamide 6 on bulk and strength of resulting BCF
   yarn)
Yarns
   (effects of online melt blending of polypropylene with
   polyamide 6 on bulk and strength of resulting BCF yarn)
Polyamides, uses
RL: PEP (Physical, engineering or chemical process); POF (Polymer in
formulation); PRP (Properties); PYP (Physical process); PROC
(Process); USES (Uses)
   (effects of online melt blending of polypropylene with
   polyamide 6 on bulk and strength of resulting BCF yarn)
Polymer blends
Polypropene fibers, properties
RL: PEP (Physical, engineering or chemical process); PRP
(Properties); PYP (Physical process); PROC (Process)
   (effects of online melt blending of polypropylene with
   polyamide 6 on bulk and strength of resulting BCF yarn)
Crystallinity
   (effects of online melt blending of polypropylene with
  polyamide 6 on property of resulting BCF yarn)
Fibers
RL: PEP (Physical, engineering or chemical process); PRP
(Properties); PYP (Physical process); PROC (Process)
   (spinning; effects of online melt blending of
  polypropylene with polyamide 6 on bulk and
   strength of resulting BCF yarn)
Microstructure
   (texture-ability; effects of online melt blending of
  polypropylene with polyamide 6 on bulk and
   strength of resulting BCF yarn)
Pressure
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(variation; effects of online melt blending of polypropylene with polyamide 6 on bulk and strength of resulting BCF yarn)

IT 108-31-6, Maleic anhydride, uses

RL: MOA (Modifier or additive use); USES (Uses)
(compatibilizer; effects of online melt blending of
polypropylene with polyamide 6 on bulk and
strength of resulting BCF yarn)

IT 25038-54-4, Polyamide 6, uses

RL: PEP (Physical, engineering or chemical process); POF (Polymer in formulation); PRP (Properties); PYP (Physical process); PROC (Process); USES (Uses)

(effects of online melt blending of polypropylene with polyamide 6 on bulk and strength of resulting BCF yarn)

IT 25085-53-4

RL: PEP (Physical, engineering or chemical process); POF (Polymer in formulation); PRP (Properties); PYP (Physical process); PROC (Process); USES (Uses)

(fibers; effects of online melt blending of polypropylene with polyamide 6 on bulk and strength of resulting BCF yarn)

IT 743436-27-3, Rolfil PN 80

RL: MOA (Modifier or additive use); PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process); USES (Uses)

(spin finish; effects of online melt blending of polypropylene with polyamide 6 on bulk and strength of resulting BCF yarn)

L133 ANSWER 10 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN

2003:801005 Document No. 140:60685 Morphology and mechanical properties of crosslinked PE/PE composite materials. Ratner, S.; Weinberg, A.; Marom, G. (Casali Institute of Applied Chemistry, The Hebrew University of Jerusalem, Jerusalem, 91904, Israel). Polymer Composites, 24(3), 422-427 (English) 2003. CODEN: PCOMDI. ISSN: 0272-8397. Publisher: Society of Plastics Engineers.

AB Novel peroxide-crosslinked composite materials based on ultra high mol. weight polyethylene fibers were studied in comparison with their non-crosslinked homologues. Two material types were prepared, either by embedding the fibers in a low-d. polyethylene matrix, or by compacting neat fibers. The resulting composites were analyzed by differential scanning calorimetry, focusing on the typical double-melting endotherm of the PE fibers, and by X-ray diffraction, concentrating on the characteristic weak 010 reflection of the triclinic phase in the fiber. Mech. testing of unidirectional composites showed crosslinking to be highly advantageous, yielding significant

property enhancement, e.g., increasing the flexural modulus from 28 to 33 GPa in the compacted fiber composite. The advantage results from the crosslinking mechanism, entailing only partial melting that is confined to the fiber skin, while retaining a significant degree of crystallinity in the fiber.

IT 9002-88-4, Polyethylene

RL: POF (Polymer in formulation); PRP (Properties); USES (Uses) (low-d.; morphol. and mech. properties of crosslinked UHMWPE/LDPE composite)

RN 9002-88-4 HCAPLUS

CN Ethene, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 74-85-1 CMF C2 H4

 $H_2C = CH_2$ 

CC 38-3 (Plastics Fabrication and Uses)

ST morphol UHMWPE fiber crosslinked LDPE composite

IT Polyolefin fibers

RL: MOA (Modifier or additive use); PRP (Properties); USES (Uses) (ethylene, Dyneema SK-75; morphol. and mech. properties of crosslinked UHMWPE/LDPE composite)

IT Mechanical properties

Polymer morphology

Young's modulus

(morphol. and mech. properties of crosslinked UHMWPE/LDPE composite)

IT 9002-88-4, Polyethylene

RL: POF (Polymer in formulation); PRP (Properties); USES (Uses) (low-d.; morphol. and mech. properties of crosslinked UHMWPE/LDPE composite)

L133 ANSWER 11 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN

2003:713624 Document No. 139:308782 Studies on the preparation of immersion-type polypropylene fragrant fiber. II.

The supermolecular structure of the matrix fiber and its sorption properties for fragrant molecules. Wang, Bing; Zhao, Jiasen (School of Material Science and Chemical Engineering, Tianjin Polytechnic University, Tianjin, 300160, Peop. Rep. China). Journal of Applied Polymer Science, 90(4), 973-981 (English) 2003. CODEN: JAPNAB. ISSN: 0021-8995. Publisher: John Wiley & Sons, Inc..

AB The supermol. structure and mech. properties of a matrix

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fiber were investigated with differential
scanning calorimetry, wide-angle X-
ray diffraction, sonic velocity instrumentation,
and fiber electronic cerimetry. The results indicated
that the apparent crystallinity of the matrix fiber, the
crystallinity of the polypropylene (PP) component,
and the mech. properties of the matrix fiber were all
affected by ethylene-vinyl acetate copolymer (EVA). Drawing
increased the orientation and tensile strength of the matrix
fiber at the proper temperature and ratio of PP to EVA.
matrix fiber, having good mech. properties, was placed in
a fragrance-alc. solution or an essential oil for the preparation of a
fragrant fiber. The fragrance property of the
fiber was determined with generally a method using slow release
time as a certain standard, and the results showed that the fragrance
property of the fiber was good.
40-4 (Textiles and Fibers)
Section cross-reference(s): 62
polypropylene fiber morphol
crystallinity sorption fragrant mol; ethylene vinyl acetate
copolymer polypropylene fiber property
Polyolefin fibers
Synthetic polymeric fibers, uses
RL: POF (Polymer in formulation); PRP (Properties); USES (Uses)
   (ethylene-vinyl acetate; supermol. structure of
   polypropylene-ethylene-vinyl acétate copolymer blend
   fiber and its sorption properties for fragrant mols.)
Adsorption
Breaking strength
Crystallinity
Elongation at break
Perfumes
Polymer morphology
Yield strength
   (supermol. structure of polypropylene-ethylene-vinyl
   acetate copolymer blend fiber and its sorption
   properties for fragrant mols.)
Polypropene fibers, uses
RL: POF (Polymer in formulation); PRP (Properties); USES (Uses)
   (supermol. structure of polypropylene-ethylene-vinyl
   acetate copolymer blend fiber and its sorption
   properties for fragrant mols.)
140-11-4, Benzyl acetate
RL: COS (Cosmetic use); PEP (Physical, engineering or chemical
process); PYP (Physical process); BIOL (Biological study); PROC
(Process); USES (Uses)
   (perfume; supermol. structure of polypropylene
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-ethylene-vinyl acetate copolymer blend fiber and its

sorption properties for fragrant mols.)

IT 24937-78-8, Ethylene-vinyl acetate copolymer 25085-53-4,

Isotactic polypropylene

RL: POF (Polymer in formulation); PRP (Properties); USES (Uses)

(supermol. structure of polypropylene-ethylene-vinyl acetate copolymer blend fiber and its sorption properties for fragrant mols.)

L133 ANSWER 12 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN

2003:425650 Document No. 140:164832 Study on in-situ fiber
reinforced composites: polyolefin/polycaprolactone blend.
Semba, T.; Kitagawa, K.; Hamada, H. (Department of Applied
Chemistry, Kyoto Municipal Institute for Industrial Research,
Shimogyo-ku, Kyoto, 600-8813, Japan). Design, Manufacturing and
Applications of Composites, Proceedings of the Joint Canada-Japan
Workshop on Composites, 4th, Vancouver, BC, Canada, Sept. 19-21,
2002, Meeting Date 2002, 176-183. Editor(s): Lo, J. CRC Press LLC:
Boca Raton, Fla. ISBN: 0-8493-1534-4 (English) 2003. CODEN: 69DYWA.

AB In-situ fiber reinforced composites were manufactured in
polymer processing process with shear and elongational flows. In

polymer processing process with shear and elongational flows. In this study, the in-situ fiber reinforced composite was prepared by blending polypropylene (PP) and polycaprolactone (PCL). This work proved that the in-situ fiber formation in PP/PCL blends were due to a shear and elongational stress. The PP/PCL sheets were manufactured by use of single screw extruder equipped with slit type die which could be applied higher shear and elongational stresses and heat press machine with lower shear and elongational stresses. The mech. properties of the extrusion sheets were considerably improved in contrast with compressive sheets attributed to the in-situ dispersed long fiber phase in extrusion processing. The DSC curves made clear the difference of crystallization between the fiber and spherical shape dispersive components.

CC 38-3 (Plastics Fabrication and Uses) Section cross-reference(s): 37

ST insitu **fiber** reinforced composite formation **polypropylene** polycaprolactone blend

IT Polyesters, processes

RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP (Physical process); PROC (Process) (caprolactone-based; in-situ fiber reinforced composites formation via blending polypropylene and polycaprolactone under shear and elongational flows and properties of their extruded and press sheets)

IT Flow

(elongational; in-situ fiber reinforced composites formation via blending polypropylene and polycaprolactone under shear and elongational flows and

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properties of their extruded and press sheets)
IT
     Crystallinity
     Fusion enthalpy
     Polymer morphology
     Stress-strain relationship
     Tensile strength
        (in-situ fiber reinforced composites formation via
        blending polypropylene and polycaprolactone under shear
        and elongational flows and properties of their extruded and press
        sheets)
IT
     Polymer blends
     Polypropene fibers, processes
     RL: PEP (Physical, engineering or chemical process); PRP
     (Properties); PYP (Physical process); PROC (Process)
       (in-situ fiber reinforced composites formation via
        blending polypropylene and polycaprolactone under shear
        and elongational flows and properties of their extruded and press
        sheets)
IT
     Flow
        (shear; in-situ fiber reinforced composites formation
        via blending polypropylene and polycaprolactone under
        shear and elongational flows and properties of their extruded and
        press sheets)
IT
     25085-53-4, Isotactic polypropylene
     RL: PEP (Physical, engineering or chemical process); PRP
     (Properties); PYP (Physical process); PROC (Process)
        (Polypro J 900GP; in-situ fiber reinforced composites
        formation via blending polypropylene and
        polycaprolactone under shear and elongational flows and
        properties of their extruded and press sheets)
IT
     24980-41-4, ε-Caprolactone homopolymer
     RL: PEP (Physical, engineering or chemical process); PRP
     (Properties); PYP (Physical process); PROC (Process)
        (assumed monomers; in-situ fiber reinforced composites
        formation via blending polypropylene and
        polycaprolactone under shear and elongational flows and
        properties of their extruded and press sheets)
IT
     25248-42-4, Celgreen PH 7
     RL: PEP (Physical, engineering or chemical process); PRP
     (Properties); PYP (Physical process); PROC (Process)
        (in-situ fiber reinforced composites formation via
        blending polypropylene and polycaprolactone under shear
        and elongational flows and properties of their extruded and press
        sheets)
L133 ANSWER 13 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
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density polyethylene fibers using online Raman

2003:156493

Document No. 138:370191 Estimating crystallinity in high

spectroscopy. Paradkar, R. P.; Sakhalkar, S. S.; He, X.; Ellison, M. S. (Center For Advanced Engineering Fibers and Films, and The School of Materials Science and Engineering, Clemson University, Clemson, SC, 29634-0971, USA). Journal of Applied Polymer Science, 88(2), 545-549 (English) 2003. CODEN: JAPNAB. ISSN: 0021-8995. Publisher: John Wiley & Sons, Inc.. AB Online Raman spectra, obtained at different points along the spin line during pilot-scale nonisothermal melt spinning of high-d. polyethylene (HDPE) fibers, are presented for the first time. The fraction of the crystalline phase corresponding to each spectrum was determined from the normalized integrated intensity of the 1418 cm-1 Raman band. It is well established that this band represents the orthorhombic crystalline phase in polyethylene. The ests. of percent crystallinity obtained from decomposition of the Raman spectrum were compared with the percent crystallinity from differential scanning calorimetry ( DSC) measurements. Online Raman spectroscopy can be successfully used to monitor the development of crystallinity in HDPE fibers as a function of distance from the spinneret. IT 9002-88-4P, Polyethylene RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (high-d., fiber; estimating crystallinity in spinning of high-d. polyethylene fibers using online Raman spectroscopy) RN9002-88-4 HCAPLUS CNEthene, homopolymer (9CI) (CA INDEX NAME) CM CRN 74-85-1 CMF C2 H4

#### $H_2C = CH_2$

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CC
     40-2 (Textiles and Fibers)
ST
     crystallinity spinning polyethylene fiber; HDPE
     fiber spinning crystallinity Raman spectroscopy
IT
     Crystallinity
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(estimating crystallinity in spinning of high-d. polyethylene fibers using online Raman spectroscopy)

IT Polyolefin fibers

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

(ethylene; estimating crystallinity in spinning of high-d.

polyethylene fibers using online Raman spectroscopy)

IT

9002-88-4P, Polyethylene

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RL: PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation)
        (high-d., fiber; estimating crystallinity in
        spinning of high-d. polyethylene fibers using online
        Raman spectroscopy)
L133 ANSWER 14 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
2002:927732
              Document No. 138:14348 Optical fiber cable
     buffer tube of dimensionally stable polyolefin.
     Castellani, Luca; Maritano, Mauro; Brandi, Giovanni (Pirelli Cavi E
     Sistemi S.P.A, Italy). PCT Int. Appl. WO 2002097503 A1 20021205, 36
          DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG,
     BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES,
     FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR,
     KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO,
     NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD,
     RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES,
     FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD,
     TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2002-EP5536
     20020521. PRIORITY: EP 2001-113193 20010530; US 2001-PV294574
     20010601.
AB
     The buffer tubes for optical fibers comprise a first
     crystalline polyolefin, i.e., propylene-alkene
     copolymers and a second crystalline polyolefin, i.e.,
     polypropylene or ethylene-propylene
     copolymer, and 0.05-1% nucleating agent,
     selected from inorg. salts, D-sorbitol derivs.,
     carboxylic acids and their salts, talc, and mixts.
     mixture of crystalline propylene-butylene copolymers
     (Clyrell 831, melt flow index (MI) 1.8 g/10 min and Moplen Ultra 925
     MI of 10 g/min and containing phenolic stabilizer), talc, and
     Millad 3988 (sorbitol derivative) nucleating
     agent was mixed at 190° and extruded to obtain tubes
     of 3.5 mm outer diameter The tubes can be extruded at low temps. and
     show good dimensional stability, no deformation, and high
     transparency, and are suitable for land and undersea deployment.
IT
     14807-96-6, Talc, uses
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); TEM (Technical or engineered material use); PROC
     (Process); USES (Uses)
        (extruded tubes of crystalline polypropylene-
        polyolefin with good dimensional stability and
        transparency for buffer casing of optical fiber cables)
RN
     14807-96-6 HCAPLUS
CN
     Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)
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#### ●3/4 Mq

Absolute stereochemistry.

IC ICM G02B006-44

CS C08L023-14; C08F210-06

CC 37-6 (Plastics Manufacture and Processing) Section cross-reference(s): 38, 73

ST polypropylene polyolefin buffer tube optical fiber cable; nucleating agent

sorbitol deriv propylene copolymer tube IT Crystal nucleating agents Extrusion of plastics and rubbers Optical cables Pipes and Tubes (extruded tubes of crystalline polypropylenepolyolefin with good dimensional stability and transparency for buffer casing of optical fiber cables) IT 29160-13-2, 1-Butene-propylene copolymer RL: PEP (Physical, engineering or chemical process); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (crystalline, Clyrell 831RCXP and Moplen Ultra 925RCXP; extruded tubes of crystalline polypropylene-polyolefin with good dimensional stability and transparency for buffer casing of optical fiber cables) 14807-96-6, Talc, uses IT RL: PEP (Physical, engineering or chemical process); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (extruded tubes of crystalline polypropylenepolyolefin with good dimensional stability and transparency for buffer casing of optical fiber cables) IT 135861-56-2, Millad 3988 RL: MOA (Modifier or additive use); PEP (Physical, engineering or chemical process); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (nucleating agent; extruded tubes of crystalline polypropylene-polyolefin with good dimensional stability and transparency for buffer casing of optical fiber cables) L133 ANSWER 15 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN Document No. 137:233957 Manufacture of polyester filaments with low modulus and high elastic

2002:727406 Document No. 137:233957 Manufacture of polyester
 filaments with low modulus and high elastic
 recovery and good winding and unwinding properties by coating the
 spun polyester filaments with lubricants to form
 filaments with specified degree of swelling. Ito, Makoto;
 Yamashita, Kenji (Nippon Ester Co., Ltd., Japan). Jpn. Kokai Tokkyo
 Koho JP 2002275731 A2 20020925, 6 pp. (Japanese). CODEN: JKXXAF.
 APPLICATION: JP 2001-80612 20010321.

AB The polyester filaments (A) are prepared by the steps
 comprising the steps of (a) melt spinning a polyester, (b) cooling
 and solidifying the fibers, and (c) coating the
 fibers with lubricants tp form fibers with a
 lubricant content to cause degree of swelling [[(E2-E1)/E1]100; E1 =

natural stretching elongation of the nonlubricated undrawn yarns in
%; E2 = natural stretching elongation of the lubricated undrawn

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yarns in %] of the fibers ≥5% and give
filaments with initial modulus ≤50 cN/
dtex and elastic recovery ≥50% at stretch 15%, or A
polyester filaments are prepared by the above steps to give
filaments with lubricant content 0.25-1.0% (on
filament). Poly(butylene terephthalate) was melt spun,
coated with an aqueous emulsion containing 50%
polyethylene-polypropylene glycol (I) with mol. wt
. (Mw) 1000 and 42% I with Mw 500, and wound to give a package with
weight kg and comprising fibers with lubricant
content 0.50% and degree of swelling 8.5% and showing unwinding
tension of the outer layer 1.5 cN, unwinding tension of the middle
layer 2.0 cN, and unwinding tension of the inner layer 3.5 cN.
spun fibers were drawn at 70° and heat-treated
110° to give yarns with tensile strength 2.9 cN/dtex
, elongation 35%, initial modulus 38 cN/dtex,
and elastic recovery 65%.
ICM D01F006-62
ICS D01F006-62
40-2 (Textiles and Fibers)
polybutylene terephthalate fiber spinning
lubricant finishing unwinding property enhancement;
polytrimethylene terephthalate fiber spinning
lubricant finishing unwinding property enhancement;
polyester fiber spinning lubricant
finishing unwinding property enhancement
Polyester fibers, uses
RL: PEP (Physical, engineering or chemical process); PRP
(Properties); PYP (Physical process); TEM (Technical or engineered
material use); PROC (Process); USES (Uses)
   (butanediol-terephthalic acid; manufacture of polyester
   filaments with low modulus and high elastic
   recovery and good winding and unwinding properties by coating the
   spun polyester filaments with lubricants to form
   filaments with specified degree of swelling)
Polyoxyalkylenes, uses
RL: MOA (Modifier or additive use); PRP (Properties); USES (Uses)
   (ethers with castor oil; manufacture of polyester filaments
   with low modulus and high elastic recovery and good
   winding and unwinding properties by coating the spun polyester
   filaments with lubricants to form filaments
   with specified degree of swelling)
Castor oil
RL: MOA (Modifier or additive use); PRP (Properties); USES (Uses)
   (ethoxylated; manufacture of polyester filaments with low
  modulus and high elastic recovery and good winding and
  unwinding properties by coating the spun polyester
   filaments with lubricants to form filaments
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with specified degree of swelling)
Hydrocarbon oils

RL: MOA (Modifier or additive use); PRP (Properties); USES (Uses) (lubricant; manufacture of polyester filaments with low modulus and high elastic recovery and good winding and unwinding properties by coating the spun polyester filaments with lubricants to form filaments with specified degree of swelling)

IT Lubricants

IT

(manufacture of polyester filaments with low modulus and high elastic recovery and good winding and unwinding properties by coating the spun polyester filaments with lubricants to form filaments with specified degree of swelling)

IT Polyester fibers, uses

RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(manufacture of polyester filaments with low modulus and high elastic recovery and good winding and unwinding properties by coating the spun polyester filaments with lubricants to form filaments with specified degree of swelling)

IT Polyester fibers, uses

RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(terephthalic acid-trimethylene glycol; manufacture of polyester filaments with low modulus and high elastic recovery and good winding and unwinding properties by coating the spun polyester filaments with lubricants to form filaments with specified degree of swelling)

IT 24968-12-5, Poly(butylene terephthalate) 26062-94-2, Poly(butylene terephthalate) 26546-03-2, Terephthalic acid-trimethylene glycol copolymer, sru 26590-75-0, Terephthalic acid-trimethylene glycol copolymer

RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(fiber; manufacture of polyester filaments with low modulus and high elastic recovery and good winding and unwinding properties by coating the spun polyester filaments with lubricants to form filaments with specified degree of swelling)

9002-92-0, Polyoxyethylene lauryl ether 9003-11-6, Polyethylenepolypropylene glycol 19149-85-0, Oleyl laurate
31565-37-4, Isotridecyl stearate

RL: MOA (Modifier or additive use); PRP (Properties); USES (Uses)

(lubricant; manufacture of polyester filaments with low modulus and high elastic recovery and good winding and unwinding properties by coating the spun polyester filaments with lubricants to form filaments with specified degree of swelling)

IT 25322-68-3D, Polyethylene glycol, ethers with castor oil RL: MOA (Modifier or additive use); PRP (Properties); USES (Uses) (manufacture of polyester filaments with low modulus and high elastic recovery and good winding and unwinding properties by coating the spun polyester filaments with lubricants to form filaments with specified degree of swelling)

L133 ANSWER 16 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
2002:503288 Document No. 137:81352 Polythiophenylene-based
gas-impermeable, high-thermal-stability polymer housing material for
fuel cell stacks. Hofmann, Achim; Radden, Peter; Reil, Frank;
Reuschel, Gerhard; Sparenberg, Bernd (Ticona G.m.b.H., Germany).
Ger. Offen. DE 10064656 A1 20020704, 8 pp. (German). CODEN:
GWXXBX. APPLICATION: DE 2000-10064656 20001222.

AB A casing material, especially for fuel cell stacks, with high thermal stability and low gas permeability coeffs., consist of a layer of liquid crystalline polymer and at least one layer of a polyarylene sulfide, preferably poly-p-phenylene sulfide. Suitable liquid crystalline polymers can include polyethylene terephthalate, poly(p-phenyleneisophthalamide), poly(m-phenyleneisophthalamide), polyester carbonates, polyazomethines, polyarylates, polythioesters, polyester polyamides, and polyester polyimides. The material, which can be used in combination with other plastics or metals, can be combined or reinforced with a number of other components, including fillers, reinforcing fibers, graft copolymers, antioxidants, nucleation agents, mold-releasing agents, etc.

IT 14807-96-6, Talc, uses

RL: DEV (Device component use); MOA (Modifier or additive use); PRP (Properties); USES (Uses)

(filler, plastic composites containing; polythiophenylene-based gas-impermeable, high-thermal-stability polymer housing material for fuel cell stacks)

RN 14807-96-6 HCAPLUS

CN Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)

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### ●3/4 Mq

CM 1

CRN 74-85-1 CMF C2 H4

#### $H_2C = CH_2$

IC ICM H01M008-02 ICS F17C001-16

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 38

IT Reinforced plastics

RL: DEV (Device component use); PRP (Properties); USES (Uses) (aramid fiber-reinforced, housing materials containing; polythiophenylene-based gas-impermeable, high-thermal-stability polymer housing material for fuel cell stacks)

IT Reinforced plastics

RL: DEV (Device component use); PRP (Properties); USES (Uses) (carbon fiber-reinforced, housing materials containing; polythiophenylene-based gas-impermeable, high-thermal-stability polymer housing material for fuel cell stacks)

IT Chalk

Mica-group minerals, uses

RL: DEV (Device component use); MOA (Modifier or additive use); PRP (Properties); USES (Uses)

(filler, plastic composites containing; polythiophenylene-based gas-impermeable, high-thermal-stability polymer housing material

for fuel cell stacks)

IT Reinforced plastics

RL: DEV (Device component use); PRP (Properties); USES (Uses) (glass fiber-reinforced, housing materials containing; polythiophenylene-based gas-impermeable, high-thermal-stability polymer housing material for fuel cell stacks)

IT Polyolefins

RL: DEV (Device component use); MOA (Modifier or additive use); PRP (Properties); USES (Uses)

(graft polymers with acrylonitrile-styrene copolymer, plastic composites containing; polythiophenylene-based gas-impermeable, high-thermal-stability polymer housing material for fuel cell stacks)

IT Silicates, uses

RL: DEV (Device component use); MOA (Modifier or additive use); PRP (Properties); USES (Uses)

(layered, filler, plastic composites containing; polythiophenylene-based gas-impermeable, high-thermal-stability polymer housing material for fuel cell stacks)

IT Reinforced plastics

RL: DEV (Device component use); PRP (Properties); USES (Uses) (mineral fiber-reinforced, housing materials containing; polythiophenylene-based gas-impermeable, high-thermal-stability polymer housing material for fuel cell stacks)

IT 471-34-1, Calcium carbonate, uses 1318-93-0, Montmorillonite, uses 7631-86-9, Silicon dioxide, uses 13983-17-0, Wollastonite 14807-96-6, Talc, uses

RL: DEV (Device component use); MOA (Modifier or additive use); PRP (Properties); USES (Uses)

(filler, plastic composites containing; polythiophenylene-based gas-impermeable, high-thermal-stability polymer housing material for fuel cell stacks)

IT 9002-84-0, PTFE 9003-54-7D, Acrylonitrile-styrene copolymer, graft copolymers with  $\alpha$  -olefins

RL: DEV (Device component use); MOA (Modifier or additive use); PRP (Properties); USES (Uses)

(plastic composites containing; polythiophenylene-based gas-impermeable, high-thermal-stability polymer housing material for fuel cell stacks)

IT 9002-88-4, Polyethylene

RL: DEV (Device component use); MOA (Modifier or additive use); PRP (Properties); USES (Uses)

(ultrahigh-d., plastic composites containing; polythiophenylene-based gas-impermeable, high-thermal-stability polymer housing material for fuel cell stacks)

L133 ANSWER 17 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN 2000:746858 Document No. 134:253567 Technological aspects of additive

use for polyolefin fiber production. Grigoriu, Aurelia; Grigoriu, George Ervant ("Gh. Asachi" Technical University, Iasi, Rom.). Plastics Engineering (New York), 59 (Handbook of Polyolefins (2nd Edition)), 825-839 (English) 2000. CODEN: PLENEZ. ISSN: 1040-2527. Publisher: Marcel Dekker, Inc.. AB A review with 115 refs. Stabilizers, flame retardants, biocides, and spin finishing additives for polyolefin fibers were introduced. Requirements of polyolefin fiber additives were discussed. CC 40-0 (Textiles and Fibers) STreview polyolefin fiber additive IT Biocides Fireproofing agents Stabilizing agents (technol. aspects of additive use for polyolefin fiber production) IT Polyolefin fibers RL: IMF (Industrial manufacture); PREP (Preparation) (technol. aspects of additive use for polyolefin fiber production) L133 ANSWER 18 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN Document No. 133:267592 Fiber-reinforced 2000:746338 polyethylene terephthalate (PET) composition. Yeum, Joon-Taek; Choe, Byung-Kil; Sonn, Young-Ho (Cheil Synthetics, Inc., S. Korea). Repub. Korea KR 132378 B1 19980413, No pp. given (Korean). KRXXFC. APPLICATION: KR 1994-18181 19940726. AB The title compns. with improved plasticity and mold release properties comprise PET having inherent viscosity of 0.5-0.9 100, inorg. filler having particle diameter 30-500 μm, e.g., mica , 0.1-40, fiber reinforcing agent, e.g., glass fiber and carbon fiber 0.1-30, inorg. nucleating agent with high dispersibility, e.g., silica, kaolin or talc 0.1-10, a polyester resin with low mol. weight or a polyolefin resin 0.5-30, and Na, Mg or Zn salt of copolymer of  $\alpha$  -olefin hydrocarbon and  $\alpha,\beta$ -unsatd. carboxylic acid 0.1-10 parts. IT 14807-96-6, Talc, uses RL: MOA (Modifier or additive use); USES (Uses) (nucleating agent; fiber-reinforced

polyethylene terephthalate composition)

Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)

RN

CN

14807-96-6 HCAPLUS

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НО- Si- ОН
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### ●3/4 Mg

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ICM C08L067-02
IC
CC
     37-6 (Plastics Manufacture and Processing)
ST
     polyethylene terephthalate fiber reinforced compn
     plasticity mold release
     Polyesters, uses
IT
     Polymer blends
     RL: POF (Polymer in formulation); TEM (Technical or engineered
     material use); USES (Uses)
        (fiber-reinforced polyethylene terephthalate composition)
IT
     Carbon fibers, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (fiber-reinforced polyethylene terephthalate composition)
IT
     Mica-group minerals, uses
     RL: MOA (Modifier or additive use); TEM (Technical or engineered
     material use); USES (Uses)
        (fillers; fiber-reinforced polyethylene terephthalate
        composition)
IT
     Polyolefins
     RL: POF (Polymer in formulation); TEM (Technical or engineered
     material use); USES (Uses)
        (low-mol. weight; fiber-reinforced polyethylene
        terephthalate composition containing)
IT
     Fillers
        (mica; fiber-reinforced polyethylene
        terephthalate composition)
ΙT
     Kaolin, uses
     RL: MOA (Modifier or additive use); USES (Uses)
        (nucleating agent; fiber-reinforced
        polyethylene terephthalate composition)
IT
     Glass fibers, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (reinforcing agents; fiber-reinforced polyethylene
        terephthalate composition)
IT
     Carboxylic acids, uses
     RL: MOA (Modifier or additive use); USES (Uses)
        (unsatd., \alpha, \beta-,
                          copolymers with
        \alpha -olefins, metal salts;
        fiber-reinforced polyethylene terephthalate composition)
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Alkenes, uses IT RL: MOA (Modifier or additive use); USES (Uses)  $(\alpha$ -, copolymers with unsatd. carboxylic acids, metal salts; fiber -reinforced polyethylene terephthalate composition) IT 25038-59-9, Poly(ethylene terephthalate), uses RL: POF (Polymer in formulation); TEM (Technical or engineered material use); USES (Uses) (fiber-reinforced polyethylene terephthalate composition) 7631-86-9, Silica, uses 14807-96-6, Talc, uses IT RL: MOA (Modifier or additive use); USES (Uses) (nucleating agent; fiber-reinforced polyethylene terephthalate composition) L133 ANSWER 19 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN 2000:615179 Document No. 133:194497 Preliminary study on domestic UHMWPE fiber. Hu, Zuming; Yu, Junrong; Liu, Zhaofeng; Lin, Jiguang (State Key Laboratory for Modification of Chemical Fibers and Polymer Materials, Donghua University, Shanghai, 200051, Peop. Rep. China). Hecheng Xianwei Gongye, 23(4), 13-15 (Chinese) 2000. CODEN: HXIGEU. ISSN: 1001-0041. Publisher: Yueyang Shiyou Huagong Zhongchang Yanjiuyuan. The structure and mech. properties of domestic UHMWPE fiber AΒ were studied by mech. property tester, acousticmeter, crystalline orientation tester, WAXD and DSC, and compared with those of imported UHMWPE fiber. The mech. properties and crystalline structure of domestic UHMWPE fiber were similar to those of imported fiber, the elongation at break was higher, and the amorphous orientation lower. The domestic fiber can be drawn at higher ratio to achieve better mech. properties. IT 9002-88-4, Polyethylene RL: PRP (Properties) (fibers; crystalline structure, mech. and thermal properties of UHMWPE fiber made in China) 9002-88-4 HCAPLUS RN CN Ethene, homopolymer (9CI) (CA INDEX NAME) CM CRN 74-85-1

 $H_2C = CH_2$ 

CC 40-4 (Textiles and Fibers)
ST UHMWPE fiber mech thermal crystallinity;

CMF C2 H4

polyethylene fiber mech ΙT Crystal structure Crystallinity Fusion enthalpy (crystalline structure, mech. and thermal properties of UHMWPE fiber made in China) IT Polyolefin fibers RL: PRP (Properties) (ethylene; crystalline structure, mech. and thermal properties of UHMWPE fiber made in China) IT 9002-88-4, Polyethylene RL: PRP (Properties) (fibers; crystalline structure, mech. and thermal properties of UHMWPE fiber made in China) L133 ANSWER 20 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN 2000:177063 Document No. 132:309630 Structural changes of gel drawn, ultra-high molecular weight polyethylene fibers with kerosene as a solvent. Xiao, C. F.; Zhang, Y. F.; An, S. L.; Jia, G. X. (Department of Materials Science, Tianjin Institute of Textile Science and Technology, Tianjin, 300160, Peop. Rep. China). Polymer Engineering and Science, 40(1), 238-246 (English) 2000. CODEN: ISSN: 0032-3888. Publisher: Society of Plastics Engineers. ABFibers of ultra-high mol. weight polyethylene (UHMW-PE) were prepared with the gel fiber drawing method. The solvent and extraction solvent used were a general kerosene and qasolinė, resp. structural changes in the fibers during the drawing process were examined by wide-angle X-ray diffraction (WAXD), polarizing optical microscopy (POM), and a d. gradient column. During the drawing process, a folded chain structure transformed into an extended-chain in the fiber, the d. of the fibers increased, and the dimensions of the orthorhombic crystallite size tended towards a narrow distribution simultaneously with the increasing draw ratio. The orthorhombic unit cell dimensions for the fiber with a draw ratio of 40 were a = 0.732 nm, b = 0.491 nm, and c = 0.254 nm. In the fiber with a draw ratio of about 40, kink bands were observed It is suggested they correspond with the local crystalline slip due to the effect of winding tension. Also, the smaller the winding roller diameter, the more clearly they can be observed In this work, the effect of the

IT 9002-88-4, Polyethylene

observed

crystallinity of fiber bulk was not

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(structural changes of gel drawn, ultra-high mol. weight

existence of kink bands on the structures of orientation and

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polyethylene fibers with kerosene as solvent)
     9002-88-4 HCAPLUS
RN
CN
     Ethene, homopolymer (9CI) (CA INDEX NAME)
     CM
     CRN
         74-85-1
     CMF
          C2 H4
H_2C = CH_2
CC
     40-4 (Textiles and Fibers)
ST
     polyethylene fiber structure crystallinity; gel
     drawn polyethylene fiber
IT
     Polyolefin fibers
     RL: PRP (Properties); TEM (Technical or engineered material use);
     USES (Uses)
        (ethylene; structural changes of gel drawn, ultra-high mol. weight
        polyethylene fibers with kerosene as solvent)
IT
     Crystallinity
     Polymer chains
        (structural changes of gel drawn, ultra-high mol. weight
        polyethylene fibers with kerosene as solvent)
IT
     Kerosene
     RL: NUU (Other use, unclassified); USES (Uses)
        (structural changes of gel drawn, ultra-high mol. weight
        polyethylene fibers with kerosene as solvent)
IT
     9002-88-4, Polyethylene
     RL: PRP (Properties); TEM (Technical or engineered material use);
     USES (Uses)
        (structural changes of gel drawn, ultra-high mol. weight
        polyethylene fibers with kerosene as solvent)
L133 ANSWER 21 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1999:631331
             Document No. 131:244556 Manufacture of synthetic
     fiber tow with reduced yarn slippage on take-up rolls for
     reduced residual waste yarn formation during the drawing step and
     good drawability. Yokoyama, Masao; Kamata, Shigeyoshi; Nakano,
     Akira (Toray Industries, Inc., Japan). Jpn. Kokai Tokkyo Koho JP
     11269716 A2 19991005 Heisei, 4 pp. (Japanese). CODEN: JKXXAF.
     APPLICATION: JP 1998-68179 19980318.
AB
     Synthetic fiber tow is prepared by melt spinning
     polymers, finishing the fibers to form a tow
     with H2O content (S) 15-35%, and taking up the tow using a take-up
     roll to cause ratio of denier (D) of the tow to width (W) of the tow
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on the take-up roll in m 1x106-3x106. A polyester was melt spun,

coated with an aqueous dispersion containing 0.3% alkyl phosphate salt to form a tow with D 15x104 denier and S 30%, taken up using a take-up roll (D/W 2.73x106), and fed to a container for a certain period. The spun tow was sep. taken up from 25 containers and drawn together to give fibers with good drawability and low amount of residual waste yarns in the container. IT 9002-88-4, Polyethylene RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PROC. (Process): USES (Uses) (fiber; manufacture of synthetic fiber tow with reduced yarn slippage on take-up rolls for reduced residual waste yarn formation during the drawing step and good drawability) RN9002-88-4 HCAPLUS CN Ethene, homopolymer. (9CI) (CA INDEX NAME) CM 1 CRN 74-85-1 CMF C2 H4  $H_2C = CH_2$ IC ICM D01D005-096 ICS D01D007-00; D01F006-06; D01F006-60; D01F006-62; D01F006-76 CC 40-2 (Textiles and Fibers) ST polyester fiber tow manuf drawing waste redn; polyamide fiber tow manuf drawing waste redn; polyethylene fiber tow manuf drawing waste redn; polypropylene fiber tow manuf drawing waste redn; fiber synthetic tow manuf drawing waste redn IT Polyolefin fibers RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (ethylene; manufacture of synthetic fiber tow with reduced yarn slippage on take-up rolls for reduced residual waste yarn formation during the drawing step and good drawability) IT Polyamide fibers, uses Polyester fibers, uses Polypropene fibers, uses Synthetic polymeric fibers, uses RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (manufacture of synthetic fiber tow with reduced yarn

slippage on take-up rolls for reduced residual waste yarn formation during the drawing step and good drawability) Synthetic polymeric fibers, uses IT RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (polyphenyls; manufacture of synthetic fiber tow with reduced yarn slippage on take-up rolls for reduced residual waste yarn formation during the drawing step and good drawability) 9002-88-4, Polyethylene 9033-83-4, Polyphenylene IT 25085-53-4, Isotactic polypropylene RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (fiber; manufacture of synthetic fiber tow with reduced yarn slippage on take-up rolls for reduced residual waste yarn formation during the drawing step and good drawability) L133 ANSWER 22 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN 1999:421143 Document No. 131:103165 Polyketone compositions for water pump connectors of automobiles with high strength and good heat and coolant resistance and dimensional stability. Yamauchi, Koji; Kanomata, Akinori; Inoue, Toshihide (Toray Industries, Inc., Japan). Jpn. Kokai Tokkyo Koho JP 11181273 A2 19990706 Heisei, 10 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1997-355827 19971224. AB Title connectors are obtained by molding compns. comprising polyketones consisting of x mol% -CH2CH2CO- and y mol% -CH2C(Me)HCO- $(y/x \ 0.01-0.10)$ , optionally fibrous and/or nonfibrous fillers, rubber powders (Tg ≤20°) dispersed in the polyketones, epoxides, and nucleating agents. Thus, 100 parts polyketone prepared from ethylene, propylene and CO was mixed with 30 parts glass fiber, and molded to give a test piece showing good Izod impact strength 80 J/m, tensile strength retention 60% after treating with 50% long life coolant solution for 500 h at 130°, and good dimensional stability. IT .14807-96-6, LMS 300, uses RL: MOA (Modifier or additive use); USES (Uses) (nucleating agent; polyketone compns. for water pump connectors of automobiles with high strength and good heat and coolant resistance and dimensional stability)

14807-96-6 HCAPLUS

Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)

RN CN О || НО- Si- ОН

#### ●3/4 Mq

IC ICM C08L073-00 ICS C08G067-02; C08K003-00; C08K003-24; C08K003-34; C08K003-36; C08K005-098; C08K005-15; C08K005-42; C08K007-02; C08K013-00; C08L021-00; C08L023-16; C08L063-00; F01P003-00; F01P005-10 CC 38-3 (Plastics Fabrication and Uses) ST polyketone automobile water pump connector strength; carbon monoxide ethylene propylene copolymer heat resistance; coolant resistance polyketone water pump connector; dimensional stability polyketone water pump connector IT Glass fibers, uses RL: MOA (Modifier or additive use); USES (Uses) (fillers; polyketone compns. for water pump connectors of automobiles with high strength and good heat and coolant resistance and dimensional stability) IT Carbonates, uses Clays, uses Kaolin, uses Mica-group minerals, uses Sulfonates

RL: MOA (Modifier or additive use); USES (Uses)
(nucleating agents; polyketone compns. for
water pump connectors of automobiles with high strength and good
heat and coolant resistance and dimensional stability)

IT Crystal nucleating agents

(polyketone compns. for water pump connectors of automobiles with high strength and good heat and coolant resistance and dimensional stability)

IT Epoxides

Polyolefin rubber

RL: MOA (Modifier or additive use); USES (Uses)
(polyketone compns. for water pump connectors of automobiles with
high strength and good heat and coolant resistance and
dimensional stability)

IT Carboxylic acids, uses

RL: MOA (Modifier or additive use); USES (Uses) (salts, nucleating agents; polyketone compns. for water pump connectors of automobiles with high strength and good heat and coolant resistance and dimensional stability)

- copolymer
  RL: DEV (Device component use); IMF (Industrial manufacture); POF
  (Polymer in formulation); PREP (Preparation); USES (Uses)
   (polyketone compns. for water pump connectors of automobiles with high strength and good heat and coolant resistance and
- dimensional stability)

  IT 26061-90-5, 2-Propenoic acid, 2-methyl-, oxiranylmethyl ester, polymer with ethene

  RL: MOA (Modifier or additive use); USES (Uses)

88995-51-1P, Carbon monoxide-ethylene-propylene

(rubber; polyketone compns. for water pump connectors of automobiles with high strength and good heat and coolant resistance and dimensional stability)

- L133 ANSWER 23 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
  1999:157067 Document No. 130:224178 Parts in optical fiber
  cables using polyolefins. Risch, Brian G.; Holder, James
  D. (Alcatel Alsthom Cie. Generale d'Electricite, Fr.). Jpn. Kokai
  Tokkyo Koho JP 11064695 A2 19990305 Heisei, 12 pp. (Japanese).
  CODEN: JKXXAF. APPLICATION: JP 1998-147510 19980528. PRIORITY: US
  1997-891978 19970710.
- AB Title cable parts, preferably (multilayer) buffer tubes, fillers rods, jackets, and slots, are made of polyolefins having melt flow index (MFI) is more than .apprx.3. Alternatively, the cable parts are made of ethylene-propylene copolymers having MFI .apprx.3-10 and containing 0.05-1% nucleating agents. The parts show improved moldability associated with improved mech. strength, compression resistance, crystallinity, and solvent resistance.
- IT 14807-96-6, Talc, uses
  RL: MOA (Modifier or additive use); USES (Uses)
   (fillers; polyolefins with high melting flow index for
   parts in optical fiber cables)
- RN 14807-96-6 HCAPLUS
  CN Tale (Mc3H2(SiO2)4) (9CI) (4

IT

CN Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)

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# ●3/4 Mg

 $H_2C = CH_2$ 

CM 1

CRN 74-85-1 CMF C2 H4

 $H_2C = CH_2$ 

CC 38-3 (Plastics Fabrication and Uses) Section cross-reference(s): 73

ST optical fiber cable part polyolefin; high melt flow index polyolefin cable; ethylene propylene copolymer nucleating agent

IT Carboxylic acids, uses

RL: MOA (Modifier or additive use); USES (Uses) (alicyclic; polyolefins with high melting flow index for parts in optical fiber cables) IT Carboxylic acids, uses RL: MOA (Modifier or additive use); USES (Uses) (aliphatic, nucleating agents; polyolefins with high melting flow index for parts in optical fiber cables) IT Carboxylic acids, uses RL: MOA (Modifier or additive use); USES (Uses) (aromatic, nucleating agents; polyolefins with high melting flow index for parts in optical fiber cables) IT Fillers (carbon black; polyolefins with high melting flow index for parts in optical fiber cables) IT Plastic foams RL: DEV (Device component use); USES (Uses) (cellular polyolefins with high melting flow index for parts in optical fiber cables) IT Carbon black Kaolin Mica-group minerals (fillers; polyolefins with high melting flow index for parts in optical fiber cables) IT **Fillers** (kaolin; polyolefins with high melting flow index for parts in optical fiber cables) IT Fillers (mica; polyolefins with high melting flow index for parts in optical fiber cables) IT Optical cables (polyolefins with high melting flow index for parts in optical fiber cables) ΙT Linear low density polyethylenes RL: DEV (Device component use); USES (Uses) (polyolefins with high melting flow index for parts in optical fiber cables) IT Polyolefins RL: DEV (Device component use); PRP (Properties); USES (Uses) (polyolefins with high melting flow index for parts in optical **fiber** cables) IT 532-32-1, Sodium benzoate 582-25-2, Potassium benzoate 822-16-2, Sodium stearate RL: MOA (Modifier or additive use); USES (Uses) (alicyclic; polyolefins with high melting flow index for parts in optical fiber cables)

471-34-1, Calcium carbonate, uses 7631-86-9, Silica, uses

IT

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14807-96-6, Talc, uses
     RL: MOA (Modifier or additive use); USES (Uses)
        (fillers; polyolefins with high melting flow index for
        parts in optical fiber cables)
IT
     74-85-1D, Ethylene, polymers with \alpha
     -olefins
     RL: DEV (Device component use); USES (Uses)
        (linear-low-d.; polyolefins with high melting flow
        index for parts in optical fiber cables)
IT
     10051-44-2, Sodium caproate
                                   14047-56-4
     RL: MOA (Modifier or additive use); USES (Uses)
        (nucleating agents; polyolefins
        with high melting flow index for parts in optical fiber
        cables)
IT
     9002-88-4, Polyethylene
                               9003-07-0, Polypropylene
     9010-79-1, Ethylene-propylene copolymer
     25085-53-4, Isotactic polypropylene
     RL: DEV (Device component use); USES (Uses)
        (polyolefins with high melting flow index for parts in
        optical fiber cables)
L133 ANSWER 24 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
             Document No. 130:169030 Manufacturing olefin
     (co)polymers with controlled molecular weight. Ushioda,
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- 1133 ANSWER 24 OF 56 HCAPLUS COPYRIGHT 2005 ACS ON STN

  1999:126936 Document No. 130:169030 Manufacturing olefin

  (co)polymers with controlled molecular weight. Ushioda,
  Tsutomu; Saito, Jun; Tsutsui, Mototake; Yasuda, Yoshitoyo; Fujita,
  Hiroyuki; Uwai, Toshihiro; Ohgi, Yoshiyuki; Adachi, Minoru;
  Morimoto, Yoshitaka; Hirose, Taketo; Kugimiya, Youichi; Taniguchi,
  Masahiko; Kuramochi, Hitoshi (Chisso Corporation, Japan). PCT Int.
  Appl. WO 9907747 Al 19990218, 88 pp. DESIGNATED STATES: W: AL, AM,
  AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES,
  FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KR, KZ, LC, LK,
  LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU,
  SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW,
  AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH,
  CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR,
  NE, NL, PT, SE, SN, TD, TG. (Japanese). CODEN: PIXXD2.
  APPLICATION: WO 1998-JP3597 19980812. PRIORITY: JP 1997-217899
  19970812; JP 1997-218756 19970813; JP 1997-251689 19970917; JP
  1997-273577 19970919.
- AB Olefin (co)polymers with controlled mol. weight from 30,000 to 10,000,000 and excellent rigidity, heat resistance and transparence, usable for various industrial parts, containers, films, sheets, fibers are manufactured by use of specific catalysts and selecting polymerization time. Thus, a catalyst 100 g from dimethylsilyl(2,3,5-trimethylcyclopentadienyl)(2',4',5'-trimethylcyclopentadienyl)zirconium dichloride 0.39 g, methylaluminoxane 267 mmol and silica 10 g was mixed with n-hexane 80 mL, trimethylaluminum 0.5 mmol and propylene was supplied and

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polymerized to give a polypropylene powder 15.5 g, showing MFR
     93 g/10 min., mol. weight 86917, Mw/Mn 2.5.
IT
     14807-96-6, Talc, uses
     RL: MOA (Modifier or additive use); USES (Uses)
        (crystal nucleating agent; manufacturing
        olefin polymers with controlled mol. weight)
     14807-96-6 HCAPLUS
RN
CN
     Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)
    0
{\tt HO-Si-OH}
●3/4 Mq
IC
    ICM
          C08F004-642
     ICS C08F010-00; C08F010-06; C08F297-08; C08L023-10; C08K003-00;
          C08K005-00; C08J005-18; B29C045-00; D01F006-04; D04H001-54
CC
     37-3 (Plastics Manufacture and Processing)
ST
     olefin copolymer controlled mol wt catalyst;
     polypropylene catalyst dimethylsilyltrimethylcyclopentadieny
     1 trimethylcyclopentadienyl zirconium dichloride methylaluminoxane
IT
     Aluminoxanes
     RL: CAT (Catalyst use); USES (Uses)
        (catalyst; manufacturing olefin polymers with
        controlled mol. weight)
IT
     Crystal nucleating agents
        (manufacturing olefin polymers with controlled mol.
        weight)
IT
     Polyolefins
     RL: IMF (Industrial manufacture); PREP (Preparation)
        (manufacturing olefin polymers with controlled mol.
        weight)
IT
     Polypropene fibers, preparation
     RL: IMF (Industrial manufacture); PREP (Preparation)
        (nonwoven fabric; manufacturing olefin
        polymers with controlled mol. weight)
IT
     Nonwoven fabrics
        (polypropylene fiber; manufacturing olefin
        polymers with controlled mol. weight)
IT
     97-93-8, Triethylaluminum, uses 2397-67-3, Triisopropylaluminum
     136019-48-2
                   161442-55-3
     RL: CAT (Catalyst use); USES (Uses)
        (catalyst; manufacturing olefin polymers with
```

controlled mol. weight) 14807-96-6, Talc, uses 25085-05-6, IT Poly(3-methyl-1-butene) 25498-06-0, Polyvinylcyclohexane 81541-12-0, 1,3:2,4-Di(p-methylbenzylidene)sorbitol 85209-91-2, Sodium 2,2'-methylene-bis-(4,6-di-tertbutylphenyl)phosphate 88266-74-4, Polyallyltrimethylsilane RL: MOA (Modifier or additive use); USES (Uses) (crystal nucleating agent; manufacturing olefin polymers with controlled mol. weight) 2212-81-9, 1,3-Bis(tert-butyl-peroxyisopropyl)benzene IT RL: MOA (Modifier or additive use); USES (Uses) (in manufacturing polypropylene with controlled mol. weight)

IT 9003-07-0P, Polypropylene

RL: IMF (Industrial manufacture); PREP (Preparation) (manufacturing olefin polymers with controlled mol. weight)

L133 ANSWER 25 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1998:535368 Document No. 129:217865 Melt spinning dull polyester
multifilaments containing titanium oxide at high speeds with good
packaging stability. Ezuka, Toshihige; Moriyama, Katsuo (Nippon
Ester Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 10219524 A2
19980818 Heisei, 5 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP
1997-38403 19970205.

AB The dull polyester filaments are prepared by melt spinning polyester multifilaments containing ≥1% Ti oxide (I), coating the fibers with aqueous emulsions containing 2.0-8.0% lubricants mainly comprising ethylene oxide-propylene oxide copolymer (II) in ≥2 steps, and taking up the fibers at ≥2500 m/min to give fibers with H2O content 10.0-16.0%. A composition containing poly(ethylene terephthalate) and 2.0% I was melt spun, coated with an emulsion containing 5.0% lubricant containing 94% II, subsequently coated with an emulsion containing 3.0% lubricant containing 94% II, taken up at 3000 m/min, and wound to give a packaging with twill collapsing amount <0.5% and comprising fibers with lubricant content 0.48% and H2O content 14.8%.

IC ICM D01F011-08

ICS D01D005-096; D01F006-62; D01F006-92

CC 40-2 (Textiles and Fibers)

ST polyester fiber dull manuf packaging stability; PET fiber dull manuf packaging stability; polyethylene polypropylene glycol lubricant finish polyester; titanium oxide polyester fiber dull manuf

IT Polyesters, uses

RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(fiber; melt spinning dull polyester multifilaments containing titanium oxide at high speeds with good packaging stability)

IT Lubricants

(finishes, containing ethylene oxide-propylene oxide copolymer; melt spinning dull polyester multifilaments containing titanium oxide at high speeds with good packaging stability)

IT Polyoxyalkylenes, uses

RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(lubricant finishes; melt spinning of dull polyester multifilaments containing titanium oxide at high speeds with good packaging stability)

IT Polyester fibers, uses

RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(melt spinning dull polyester multifilaments containing titanium oxide at high speeds with good packaging stability)

IT 25038-59-9, Poly(ethylene terephthalate), uses

RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(fiber; melt spinning dull polyester multifilaments containing titanium oxide at high speeds with good packaging stability)

IT 9003-11-6, Ethylene oxide-propylene oxide

copolymer
RL: PEP (Physical, engineering or chemical process); PRP

(Properties); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(lubricant finish; melt spinning of dull polyester multifilaments containing titanium oxide at high speeds with good packaging stability)

L133 ANSWER 26 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN

1998:316405 Document No. 129:29033 Structure model of oriented copolymerized HDPE monofilaments by die drawing process. Yi, Xiaosu; Zou, Xiangping; Tan, Hongsheng (Dep. Polymers, Zhejiang Univ., Hangzhou, 310027, Peop. Rep. China). Gaofenzi Xuebao (2), 227-231 (Chinese) 1998. CODEN: GAXUE9. ISSN: 1000-3304. Publisher: Kexue Chubanshe.

AB Self-reinforced HDPE monofilament were prepared by die drawing process. According to Cox model for **fiber** composite and Gibson & Ward crystalline bridges model, the reinforced mechanism was discussed to reveal that the mech. improvement of monofilament was

due to the growth of needle-like crystal with larger aspect ratio or the increase in volume fraction of the crystalline bridge. Based on the exptl. results obtained by DMA, DSC and SEM, the theor. prediction was also analyzed and discussed, and the theor. values could well-explain the monofilament behavior. 9002-88-4, Polyethylene

PI. PPD (Properties)

RL: PRP (Properties)

(high-d., DGDB 2480; orientation, crystallinity, and morphol. of oriented HDPE monofilaments from die drawing process)

RN 9002-88-4 HCAPLUS

CN Ethene, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 74-85-1 CMF C2 H4

 $H_2C = CH_2$ 

IT

CC 40-4 (Textiles and Fibers)

IT Polyolefin fibers

RL: PRP (Properties)

(ethylene; orientation, crystallinity, and morphol. of oriented HDPE monofilaments from die drawing process)

IT Crystallinity

Filaments

Polymer morphology

(orientation, crystallinity, and morphol. of oriented HDPE monofilaments from die drawing process)

IT 9002-88-4, Polyethylene

RL: PRP (Properties)

(high-d., DGDB 2480; orientation, crystallinity, and morphol. of oriented HDPE monofilaments from die drawing process)

L133 ANSWER 27 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN

1998:268573 Document No. 129:5609 Thermobondable polyolefin fibers and their production method. Stengaard, Flemming Faurby; Balslev, Henrik; Carstensen, Peter (Fibervisions A/s, Den.; Stengaard, Flemming Faurby; Balslev, Henrik; Carstensen, Peter).

PCT Int. Appl. WO 9817746 Al 19980430, 37 pp. DESIGNATED STATES: W: AL, AM, AT, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, CZ, DE, DE, DK, DK, EE, ES, FI, FI, GB, GE, GH, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, DE, DK, ES, FI, FR, GA, GB, GR,

IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1997-DK467 19971023. PRIORITY: DK 1996-1178 19961024; US 1996-29913 19961104; DK 1997-71 19970120. A method for producing hydrophobic polyolefin-containing AB fibers or filaments, in particular cardable staple fibers, comprises treating fibers with hydrophobic spin finishes after spinning and stretching, that comprise ≥1 water-insol. ester of a mono-, di-, tri- or tetrahydric alc. with mol. weight ≤500 and a branched or straight chain C12-30 fatty acid, e.g. a water-insol. ester of ethylene or propylene glycol, glycerol, neopentyl glycol, trimethylolethane or trimethylolpropane and ≥1 C12-24 (un) saturated fatty acid residue, an anionic or nonionic antistatic agent preferably being applied after crimping. Fibers produced by the method and nonwovens produced from such fibers are included. 9002-88-4P IT RL: IMF (Industrial manufacture); PEP (Physical, engineering or chemical process); PRP (Properties); PREP (Preparation); PROC (Process) (high-d., bicomponent core-sheath fibers; manufacture of

RN 9002-88-4 HCAPLUS

CN Ethene, homopolymer (9CI) (CA INDEX NAME)

fibers and nonwovens)

hydrophobic thermobondable polyolefin staple

CM 1

CRN 74-85-1 CMF C2 H4

 $H_2C = CH_2$ 

IC ICM C10M101-00

ICS D06M013-292; D06M013-224

CC 40-10 (Textiles and Fibers)

ST hydrophobic polyolefin fiber filament; cardable staple polyolefin fiber; antistatic agent polyolefin fiber; bicomponent HDPE polypropylene fiber; ester spin finish polyolefin fiber; spunbond meltblown polyolefin nonwoven

IT Alcohols, uses

RL: TEM (Technical or engineered material use); USES (Uses) (ethoxylated, spin finishes; manufacture of hydrophobic thermobondable polyolefin staple

```
fibers and nonwovens)
IT
     Polyolefin fibers
     RL: IMF (Industrial manufacture); PEP (Physical, engineering or
     chemical process); PRP (Properties); PREP (Preparation); PROC
        (ethylene, bicomponent core-sheath; manufacture of hydrophobic
        thermobondable polyolefin staple fibers and
        nonwovens)
IT
     Polyolefin fibers
     Polypropene fibers, preparation
     RL: IMF (Industrial manufacture); PEP (Physical, engineering or
     chemical process); PRP (Properties); PREP (Preparation); PROC
     (Process)
        (ethylene-propene; manufacture of hydrophobic thermobondable
        polyolefin staple fibers and nonwovens
IT
     Polysiloxanes, uses
     Waxes
     RL: MOA (Modifier or additive use); USES (Uses)
        (friction-reducing additive; manufacture of hydrophobic thermobondable
        polyolefin staple fibers and nonwovens
IT
     Esters, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (hydrophobic spin finishes; manufacture of
        hydrophobic thermobondable polyolefin staple
        fibers and nonwovens)
IT
     Nonwoven fabrics
     Waterproofing
     Wetting agents
        (manufacture of hydrophobic thermobondable polyolefin staple
        fibers and nonwovens)
IT
     Polyolefin fibers
     Polypropene fibers, preparation
     RL: IMF (Industrial manufacture); PEP (Physical, engineering or
     chemical process); PRP (Properties); PREP (Preparation); PROC
        (manufacture of hydrophobic thermobondable polyolefin staple
        fibers and nonwovens)
                  53126-06-0, Potassium butyl phosphate
IT
     53026-21-4
     RL: MOA (Modifier or additive use); USES (Uses)
        (antistatic agents; manufacture of hydrophobic thermobondable
        polyolefin staple fibers and nonwovens
IT
     9010-79-1P, Ethylene-propylene copolymer
     25085-53-4P, Isotactic polypropylene
    RL: IMF (Industrial manufacture); PEP (Physical, engineering or
```

chemical process); PRP (Properties); PREP (Preparation); PROC

```
(Process)
        (fibers; manufacture of hydrophobic thermobondable
        polyolefin staple fibers and nonwovens
IT
     9002-88-4P
     RL: IMF (Industrial manufacture); PEP (Physical, engineering or
     chemical process); PRP (Properties); PREP (Preparation); PROC
     (Process)
        (high-d., bicomponent core-sheath fibers; manufacture of
        hydrophobic thermobondable polyolefin staple
        fibers and nonwovens)
     56-81-5D, 1,2,3-Propanetriol, esters, uses
IT.
                                                  57-55-6D,
     1,2-Propanediol, esters, uses 77-85-0D, Trimethylolethane, esters
     77-99-6D, esters 107-21-1D, 1,2-Ethanediol, esters, uses
     126-30-7D, esters
                         25618-55-7D, Polyglycerol, esters
     RL: TEM (Technical or engineered material use); USES (Uses)
        (hydrophobic spin finishes; manufacture of
        hydrophobic thermobondable polyolefin staple
        fibers and nonwovens)
L133 ANSWER 28 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
             Document No. 128:35371 Hydroperoxide-initiated grafting of
1997:754565
     poly(styrene-stat-acrylonitrile) onto ultra-high modulus
     polyethylene fibers. Arnold, J. J.; Zamora, M. P.;
     Batich, C. D.; Brennan, A. B. (Department Materials Science
     Engineering, University Florida, Gainesville, FL, 32611, USA).
     Journal of Adhesion Science and Technology, 11(10), 1343-1358
     (English) 1997. CODEN: JATEE8. ISSN: 0169-4243. Publisher: VSP
     BV.
AB
     There is evidence from modeling expts. that designed interphases,
     e.g. varying graft mol. weight and/or an adjustable interphase
    modulus, will provide a method of tailoring the mech.
     response of a composite. There is no exptl. evidence yet to
     substantiate these claims in fiber-reinforced composites.
     Hence, a grafting method that allows better design of the
     interphase, i.e. an intelligent interphase, is described.
     authors' approach is to engineer intelligent interphases using a
    hydroperoxide-initiated grafting technique that provides a novel
     method of grafting by a free radical-type polymerization. This study
     describes the hydroperoxidn. and grafting of poly(styrene-stat-
     acrylonitrile) (SAN) onto ultra-high modulus polyethylene
     (UHMPE) fibers. Differential scanning
     calorimetry and fiber tensile tests were used to
    monitor the effect of the surface treatment on bulk
     crystallinity and fiber tensile properties.
     characterization methods showed that the hydroperoxidn. treatment
```

properties. The N in poly(styrene-stat-acrylonitrile) was used as a

was not deleterious to the fiber bulk

marker for ESCA in order to verify grafting. ESCA anal. also demonstrated that the grafting reaction was a surface phenomenon, not a bulk treatment. Tapping-mode atomic force microscopy (AFM) was used to image the surface morphol. of the UHMPE fibers and images of the SAN-grafted fibers revealed a distinct topog. comprised of nodules, 3-12 nm high and 10-25 nm apart. A simple hard-sphere model is presented to explain the polymer graft conformation based on the surface morphol. as imaged by AFM. The model corroborates the conclusions that the hydroperoxidn. grafting yields linear chains tethered at the amorphous surface domains.

IT 9002-88-4DP, Polyethylene, peroxidized RL: PRP (Properties); SPN (Synthetic preparation); PREP

(Preparation)

(fibers; grafting of poly(styrene-acrylonitrile) onto hydroperoxidized ultra-high modulus polyethylene fibers)

RN 9002-88-4 HCAPLUS

CN Ethene, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 74-85-1 CMF C2 H4

 $H_2C = CH_2$ 

- CC 37-4 (Plastics Manufacture and Processing)
- ST polyethylene **fiber** hydroperoxidized grafting polystyrene acrylonitrile
- IT Polyolefin fibers

RL: PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process)

(ethylene; grafting of poly(styrene-acrylonitrile) onto hydroperoxidized ultra-high modulus polyethylene fibers)

IT Polymer morphology

(surface; grafting of poly(styrene-acrylonitrile) onto hydroperoxidized ultra-high modulus polyethylene fibers)

IT 9002-88-4DP, Polyethylene, peroxidized 25087-32-5P, Acrylonitrile-ethylene-styrene copolymer

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

(fibers; grafting of poly(styrene-acrylonitrile) onto hydroperoxidized ultra-high modulus polyethylene

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fibers)
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(grafting of poly(styrene-acrylonitrile) onto hydroperoxidized ultra-high modulus polyethylene fibers)

L133 ANSWER 29 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN

1997:618917 Document No. 127:279066 Poly(ethylene terephthalate)-based polyester compositions with improved mechanical strength and moisture-heat resistance. Matsumoto, Kazuaki (Kanegafuchi Chemical Industry Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 09241490 A2 19970916 Heisei, 12 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1996-85813 19960313.

AB Title compns. contain (A) 100 parts poly(ethylene terephthalate) - based polyesters prepared by polymerization using Ge compound catalysts, (B) 0.2-50 parts graft copolymers obtained by contacting (a) 100 parts polyolefins with aqueous suspensions containing (b) 0.1-500 parts vinyl monomers, (c) 0.1-30 parts CH2:CRC(:O)NHCH2Ar (Ar = ≥1 glycidyloxy-containing C6-23 aromatic hydrocarbyl; R = H, Me), and (d) 0.01-10 parts [for 100 parts of (a + b)] radically polymerization initiators, and (C) 5-200 parts reinforcement fillers. Thus, poly(ethylene terephthalate) prepared using GeO2 100, N-[(2,3-epoxypropoxy)-3,5dimethylphenylmethyl]acrylamide-ethylene-propylene-styrene graft copolymer 10, T 195H/P (glass fiber) 50, ADK Stab AO 60 0.35, and ADK Stab AO 412S 0.15 part were dry blended, melt kneaded, pelletized, and injection molded to give a test piece showing tensile strength 150 MPa and good moisture-heat resistance.

IT 14807-96-6, Micro Ace K 1, uses

RL: MOA (Modifier or additive use); USES (Uses) (crystal nucleating agents; poly(ethylene terephthalate)-based polyester blends with improved mech. strength and moisture-heat resistance)

RN 14807-96-6 HCAPLUS

CN Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)

|| но-si-он

●3/4 Mg

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IC
     ICM C08L067-02
     ICS C08K003-00; C08K005-00; C08G063-85; C08L067-02; C08L051-06
CC
     37-6 (Plastics Manufacture and Processing)
     Section cross-reference(s): 35, 38
ST
     polyester grafted polyolefin blend mech strength;
     polyethylene terephthalate polyolefin blend mech strength;
     glycidyl polyolefin grafted blend polyester; moisture
     resistance polyester grafted polyolefin; heat resistance
     polyester grafted polyolefin; germanium polymn catalyst
     polyethylene terephthalate
IT
     Mica-group minerals, uses
     RL: MOA (Modifier or additive use); USES (Uses)
        (crystal nucleating agents, A 21S;
        poly(ethylene terephthalate)-based polyester blends with improved
        mech. strength and moisture-heat resistance)
IT
     Reinforced plastics
     RL: PRP (Properties); TEM (Technical or engineered material use);
     USES (Uses)
        (fiber-reinforced; poly(ethylene terephthalate)-based
        polyester blends with improved mech. strength and moisture-heat
        resistance)
IT
     Polyolefins
       Polyolefins
     RL: MOA (Modifier or additive use); USES (Uses)
        (graft; poly(ethylene terephthalate)-based polyester blends with
        improved mech. strength and moisture-heat resistance)
IT
     Glass fibers, uses
     RL: MOA (Modifier or additive use); USES (Uses)
        (reinforcements, T 195H/P; poly(ethylene terephthalate)-based
        polyester blends with improved mech. strength and moisture-heat
        resistance)
IT
     14807-96-6, Micro Ace K 1, uses
     RL: MOA (Modifier or additive use); USES (Uses)
        (crystal nucleating agents; poly(ethylene
        terephthalate) - based polyester blends with improved mech.
        strength and moisture-heat resistance)
IT
     106-89-8, Epichlorohydrin, reactions
                                            13579-23-2,
     4-Acrylamidomethyl-2,6-dimethylphenol
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (preparation of modifiers for grafted polyolefins)
L133 ANSWER 30 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1996:722186
             Document No. 125:331503 Rewettable polyolefin
     fibers for nonwoven fabric layer in diapers.
    Carstensen, Peter; Revsbaaek, Per; Dyrmose-Jensen, Katharine
     (Danaklon A/s, Den.; Carstensen, Peter; Revsbaaek, Per;
    Dyrmose-Jensen, Katharine). PCT Int. Appl. WO 9633303 Al 19961024,
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38 pp. DESIGNATED STATES: W: AU, BR, CA, CN, CZ, CZ, DE, DK, EE,

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EE, GE, HU, IS, JP, KR, LT, LV, MX, NO, NZ, PL, RU, SG, SI, SK, SK,
     UA, US, VN; RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU,
     MC, NL, PT, SE. (English). CODEN: PIXXD2. APPLICATION: WO
     1996-DK178 19960419. PRIORITY: DK 1995-468 19950421.
AB
     Synthetic fibers are prepared by applying to spun
     filaments a first spin finish comprising
    ≥1 hydrophilic lubricant, stretching the filaments,
     applying to the stretched filaments a second spin
     finish comprising ≥1 cationic antistatic agent, and
     crimping, drying, and cutting the filaments to obtain
    hydrophilic staple fibers.
                                The spin
     finishes, which may each contain a hydrophilic lubricant and
     a cationic antistatic agent, may also contain a small amount of a
    polydiorganosiloxane. The fibers may be carded at high
     speeds and are useful for the preparation of hydrophilic nonwoven
    materials that can maintain wettability after one or several
    wettings. Polypropylene fibers were
     finished (both spin finishes of fatty
     alc. ethoxylate lubricant and fatty acid/polyamine condensate
     antistat), crimped, annealed, cut to staple fibers of
     fineness 2.2 dtex, carded, and thermally bonded
     (161°) to form nonwovens having tensile strength
     (machine direction) 42.0 g, bondability index 23, and rewet 0.3%,
    vs. 36.3, 16.1, and 0.26, resp., using anionic ethoxylated silicone
     spin finish.
IC
     ICM D01F006-04
     ICS D06M013-46; D06M015-647
ICI
    D06M101-20
CC
    40-9 (Textiles and Fibers)
ST
    polyolefin fiber hydrophilic finish
    nonwoven; diaper water repellent nonwoven
    finished; rewettable hydrophilic finished polyolefin
     fiber
IT
    Quaternary ammonium compounds, uses
    RL: PRP (Properties); TEM (Technical or engineered material use);
    USES (Uses)
        (finish for polyolefin fibers; rewettable
       hydrophilic polyolefin fibers for
       nonwoven fabric layer in diapers)
IT
    Lubricants
        (hydrophilic, in finish for polyolefin fibers
        ; rewettable hydrophilic polyolefin fibers
        for nonwoven fabric layer in diapers)
IT
    Antistatic agents
        (in finish for polyolefin fibers; rewettable
       hydrophilic polyolefin fibers for
       nonwoven fabric layer in diapers)
IT
    Siloxanes and Silicones, uses
```

```
RL: PRP (Properties); TEM (Technical or engineered material use);
     USES (Uses)
        (in finish for polyolefin fibers; rewettable
        hydrophilic polyolefin fibers for
        nonwoven fabric layer in diapers)
IT
     Diapers
        (rewettable hydrophilic polyolefin fibers for
        nonwoven fabric layer in)
IT
     Textile easy-care finishing
        (rewettable hydrophilic polyolefin fibers for
        nonwoven fabric layer in diapers)
IT
     Polypropene fibers, uses
     RL: PEP (Physical, engineering or chemical process); PRP
     (Properties); TEM (Technical or engineered material use); PROC
     (Process); USES (Uses)
        (rewettable hydrophilic polyolefin fibers for
        nonwoven fabric layer in diapers)
ΙT
     Alcohols, uses
     Amides, uses
     Esters, uses
     RL: PRP (Properties); TEM (Technical or engineered material use);
     USES (Uses)
        (alkoxy, lubricant for polyolefin fibers;
        rewettable hydrophilic polyolefin fibers for
        nonwoven fabric layer in diapers)
IT
     Alcohols
     RL: PRP (Properties); TEM (Technical or engineered material use);
     USES (Uses)
        (ethoxylated, finish for polyolefin fibers;
        rewettable hydrophilic polyolefin fibers for
        nonwoven fabric layer in diapers)
IT
     Textiles
        (nonwoven, rewettable hydrophilic polyolefin
        fibers for nonwoven fabric layer in diapers)
IT
     25085-53-4, Isotactic polypropylene
     RL: PEP (Physical, engineering or chemical process); PRP
     (Properties); TEM (Technical or engineered material use); PROC
     (Process); USES (Uses)
        (fiber; rewettable hydrophilic polyolefin
        fibers for nonwoven fabric layer in diapers)
L133 ANSWER 31 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
             Document No. 125:95452 Microporous membranes containing
    polyolefins and ion-exchange fibers. Nago, Kunya (Tokuyama
    Corp, Japan). Jpn. Kokai Tokkyo Koho JP 08117576 A2 19960514
    Heisei, 11 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP
    1994-257021 19941021.
AB
    The microporous membranes comprise (a) 15-70 weight%
```

polyolefins, (b) 84-20 weight% filler particles (average particle size  $0.01-5.0 \mu m$ ) dispersed in the polyolefins , and (c) 1-10 weight% ion-exchange fibers (fiber diameter ≤30 μm, average fiber length 200-2000 μm) dispersed in the polyolefins in parallel with the membrane surfaces and in approx. the same directions as to the fiber axes, the membranes having network structures comprising continuous pores with maximum pore size ≤3.0 μm, showing void volume 20-90%, and the polyolefins being mol. oriented at the direction of fiber axes of the ion-exchange fibers. The membranes provide efficient removal of contaminants (e.g. trihalomethanes, microorganisms, and other particles) in water and are also useful for air filters, wastewater treatment, etc. 14807-96-6, Talc, uses

IΤ

RL: MOA (Modifier or additive use); TEM (Technical or engineered material use); USES (Uses)

(microporous membranes containing polyolefins, filler particles, and ion-exchange fibers for purification of water and air) 14807-96-6 HCAPLUS

CN Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)

HO-Si-OH

RN

## ●3/4 Mg

RN

IT 9002-88-4, Hizex 1300J

RL: POF (Polymer in formulation); TEM (Technical or engineered material use); USES (Uses)

(microporous membranes containing polyolefins, filler particles, and ion-exchange fibers for purification of water and air) 9002-88-4 HCAPLUS

CN Ethene, homopolymer (9CI) (CA INDEX NAME)

CM

CRN 74-85-1 CMF C2 H4

 $H_2C = CH_2$ 

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IC
     ICM B01D071-26
     ICS B01D039-16; B01D071-82; C08J009-00
ICI
     C08L023-02
     61-5 (Water)
CC
     Section cross-reference(s): 59, 60
     porous membrane polyolefin ion exchanger; water purifn
ST
    porous membrane polyolefin; fiber ion exchanger water
    purifn; air filter polyolefin ion exchanger
IT
     Siloxanes and Silicones, uses
     RL: MOA (Modifier or additive use); TEM (Technical or engineered
     material use); USES (Uses)
        (Trefil R 925; microporous membranes containing polyolefins
        , filler particles, and ion-exchange fibers for purification of water
        and air)
IT
     Synthetic fibers, polymeric
     RL: POF (Polymer in formulation); TEM (Technical or engineered
    material use); USES (Uses)
        (divinylbenzene-styrene; microporous membranes containing
       polyolefins, filler particles, and ion-exchange fibers
        for purification of water and air)
IT
     Cation exchangers
     Ion exchangers
        (microporous membranes containing polyolefins, filler
        particles, and ion-exchange fibers for purification of water and air)
IT
     Silsesquioxanes
    RL: MOA (Modifier or additive use); TEM (Technical or engineered
     material use); USES (Uses)
        (Me, microporous membranes containing polyolefins, filler
        particles, and ion-exchange fibers for purification of water and air)
IT
    Air purification
    Wastewater treatment
    Water purification
        (filtration, microporous membranes containing polyolefins,
        filler particles, and ion-exchange fibers for purification of water
IT
     9003-70-7, Divinylbenzene-styrene copolymer
    RL: POF (Polymer in formulation); TEM (Technical or engineered
    material use); USES (Uses)
        (fiber; microporous membranes containing polyolefins,
        filler particles, and ion-exchange fibers for purification of water
       and air)
IT
     471-34-1, Tunex E, uses
                              9017-37-2, MP 3000 14807-96-6,
    Talc, uses
                  175069-63-3, KT 115
    RL: MOA (Modifier or additive use); TEM (Technical or engineered
    material use); USES (Uses)
        (microporous membranes containing polyolefins, filler
       particles, and ion-exchange fibers for purification of water and air)
    9002-88-4, Hizex 1300J 9003-07-0, PN 120 9010-79-1, MS
IT
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L133 ANSWER 32 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN

1995:931282 Document No. 123:343271 Cardable, hydrophobic

polyolefin staple fibers, their manufacture, and
nonwoven webs from. Jensen, Arne; Dyrmose-Jensen,
Katharine; Clausen, Lydia Dahl; Marcher, Bjoern (Danaklon A/S,
Den.). PCT Int. Appl. WO 9519465 A1 19950720, 56 pp. DESIGNATED
STATES: W: AU, BR, CA, CN, CZ, DE, DK, JP, KR, MX, RU, UA, US; RW:
AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE.
(English). CODEN: PIXXD2. APPLICATION: WO 1995-DK24 19950113.
PRIORITY: DK 1994-70 19940114.

AB Title manufacture comprises applying to spun filaments a first spin finish comprising ≥1 cationic antistatic agent, especially a quaternary ammonium salt, stretching the filaments, applying a second spin finish dispersion comprising ≥1 hydrophobic lubricant selected from a fatty acid amide condensation product and a hydrocarbon wax, crimping, drying, and cutting the filaments to obtain staple fibers. The fibers can be carded at extremely high speeds and are particularly suitable for use in the preparation of thermally bonded hydrophobic nonwoven fabrics in which a dry, water-repellent surface which can function as a liquid barrier is desired, e.g., for disposable diapers, feminine hygienic products and medical products. The second finish may further comprise a cationic antistatic agent in ≤20 weight%.

IC ICM D06M013-02

ICS D06M015-227; D06M013-463; D06M013-402

CC 40-7 (Textiles and Fibers)

ST cardable hydrophobic polyolefin staple fiber finish; fatty amide cardable hydrophobic polyolefin staple; antistatic agent cardable hydrophobic polyolefin staple; quaternary cardable hydrophobic polyolefin staple

IT Quaternary ammonium compounds, uses

RL: TEM (Technical or engineered material use); USES (Uses) (antistatic agents; in production of cardable, hydrophobic polyolefin staple fibers)

IT Paraffin waxes and Hydrocarbon waxes, uses
RL: TEM (Technical or engineered material use); USES (Uses)

(in production of cardable, hydrophobic polyolefin staple fibers)

- IT Polyolefin fibers
  - Polypropene fibers, processes
  - RL: PEP (Physical, engineering or chemical process); PROC (Process) (manufacture of cardable, hydrophobic staple)
- IT Antistatic agents
  - (quaternary ammonium compds.; in production of cardable, hydrophobic polyolefin staple fibers)
- IT Amides, uses
  - RL: TEM (Technical or engineered material use); USES (Uses) (fatty, in production of cardable, hydrophobic polyolefin staple fibers)
- IT 25085-53-4, Isotactic polypropylene
  - RL: PEP (Physical, engineering or chemical process); PROC (Process) (fiber; manufacture of cardable, hydrophobic staple)
- 9016-00-6, Dimethylsilanediol homopolymer, sru 31900-57-9, Dimethylsilanediol homopolymer RL: TEM (Technical or engineered material use); USES (Uses)
  - (in production of cardable, hydrophobic polyolefin staple fibers)
- L133 ANSWER 33 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
  1995:918311 Document No. 124:90097 Evaluation of the degree of
  crystallinity of fibers from polymer blends. Part
  I. Kwiatkowski, Ryszard; Wlochowicz, Andrzej; Rabiej, Stanislaw
  (Inst. Wlokienniczy, Politech. Lodzka, Bielsko-Biala, Pol.).
  Przeglad Wlokienniczy + Technik Wlokienniczy, 49(8), 8-10 (Polish)
  1995. CODEN: PWTWEA. ISSN: 1230-0381. Publisher: Wydawnictwo
  SIGMA NOT.
- AB A method was proposed for differentiating in the xray diffraction patterns of fibers
  prepared from a blend of a semicryst. polymer with an amorphous
  polymer (S-A samples) or from a blend of a semicryst. polymer with a
  an amorphous copolymer (C-CA samples) between the diffraction of the
  amorphous and the semicryst. portion of the fiber. The
  method was applied with good results to characterization of
  fibers prepared from polypropylene-polystyrene,
  polyethylene-ABS polymer, poly(butylene terephthalate) (I)-nylon 6,

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and I-nylon 11 blends.
CC
     40-4 (Textiles and Fibers)
     Section cross-reference(s): 37
     synthetic fiber blend x ray
ST
     diffraction; crystallinity synthetic fiber
     polymer blend; nylon 11 blend synthetic fiber
     crystallinity; polybutylene terephthalate synthetic
     fiber crystallinity; polycaprolactam blend
     synthetic fiber crystallinity; ABS polymer blend
     synthetic fiber crystallinity; polyethylene
     blend synthetic fiber crystallinity; polystyrene
     blend synthetic fiber crystallinity
IT
     Crystallinity
        (evaluation of the degree of crystallinity of
        fibers from polymer blends)
IT
     Synthetic fibers, polymeric
     RL: PRP (Properties); TEM (Technical or engineered material use);
     USES (Uses)
        (evaluation of the degree of crystallinity of
        fibers from polymer blends)
ΙT
     Polyamide fibers, uses
     RL: PRP (Properties); TEM (Technical or engineered material use);
     USES (Uses)
        (6, evaluation of the degree of crystallinity of
        fibers from polymer blends)
ΙT
     Acrylic fibers, uses
     Synthetic fibers, polymeric
     RL: PRP (Properties); TEM (Technical or engineered material use);
     USES (Uses)
        (acrylonitrile-butadiene-styrene, evaluation of the degree of
        crystallinity of fibers from polymer blends)
IT
     Polyester fibers, uses
     RL: PRP (Properties); TEM (Technical or engineered material use);
     USES (Uses)
        (butanediol-terephthalic acid, evaluation of the degree of
        crystallinity of fibers from polymer blends)
IT
     Polyolefin fibers
     RL: PRP (Properties); TEM (Technical or engineered material use);
    USES (Uses)
        (ethylene, evaluation of the degree of crystallinity of
        fibers from polymer blends)
IT
     Polyamide fibers, uses
    RL: PRP (Properties); TEM (Technical or engineered material use);
    USES (Uses)
        (laurolactam, evaluation of the degree of crystallinity
        of fibers from polymer blends)
IT
     Synthetic fibers, polymeric
    RL: PRP (Properties); TEM (Technical or engineered material use);
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USES (Uses)
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(styrene, evaluation of the degree of **crystallinity** of **fibers** from polymer blends)

IT 9003-56-9, ABS polymer 24968-12-5, Butanediol-terephthalic acid copolymer SRU 25035-04-5, Nylon 11 25587-80-8 26062-94-2, Butanediol-terephthalic acid copolymer RL: PRP (Properties): TEM (Technical or engineered material use):

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(evaluation of the degree of crystallinity of fibers from polymer blends)

L133 ANSWER 34 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1995:703798 Document No. 123:85785 Morphology of gel-spun polyethylene
fibers. Khosravi, Nasrin; Warner, S. B.; Murthy, N. S.;
Kumar, Satish (Sch. Textile and Fiber Engineering, Georgia Inst.
Technology, Atlanta, GA, 30332-0295, USA). Journal of Applied
Polymer Science, 57(7), 781-7 (English) 1995. CODEN: JAPNAB. ISSN:
0021-8995. Publisher: Wiley.

AB HNO3 etching studies were conducted on samples of com. available highly oriented gel-spun polyethylene fibers Spectra 900 and 1000 from Allied Signal, Inc. The results show that the acid attacks the fiber, increases crystallinity as observed by X-ray diffraction, increases the enthalpy of melting of unconstrained fibers by 13-25%, and removes topol. constraints, thus facilitating the crystallization of chain segments in the noncryst. regions. The acid functionalizes the fiber, creating various O- and N-containing moieties, specifically (C=O)-, -C=C-, and -NO2 groups. The small weight loss upon etching, less than 2%, and the fact that fibers weaken but do not fragments suggests that the gel-spun and -drawn morphol. is more resistant to acid attack than are other morphologies of polyethylene. Photomicroscopy shows that acid etching opens cracks normal to the fiber axis. That the acid can attack the gel-spun fibers indicates the presence of structural imperfections such as folds, mol. kinds, or uncrystd. regions within the fibers. IR anal. on virgin fibers shows the absence of absorption bands normally associated with the presence of chains lying within amorphous regions in polyethylene. These and other morphol. features are integrated into a description of the structure of com. gel-spun polyethylene fibers.

IT 9002-88-4, Polyethylene

RL: PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process)

(fibers, Spectra 900 and Spectra 1000; morphol. of acid-etched gel-spun polyethylene fibers)

RN 9002-88-4 HCAPLUS

CN Ethene, homopolymer (9CI) (CA INDEX NAME)

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CM 1
CRN 74-85-1
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C2 H4

 $H_2C = CH_2$ 

CMF

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CC 40-4 (Textiles and Fibers)
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ST gel spun polyethylene fiber morphol

IT Polyolefin fibers

RL: PEP (Physical, engineering or chemical process); PRP

(Properties); PROC (Process)

(ethylene, Spectra 900 and Spectra 1000; morphol. of acid-etched gel-spun polyethylene fibers)

IT Polymer morphology

(surface, morphol. of acid-etched gel-spun polyethylene fibers)

IT 9002-88-4, Polyethylene

RL: PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process)

(fibers, Spectra 900 and Spectra 1000; morphol. of acid-etched gel-spun polyethylene fibers)

IT 7697-37-2, Nitric acid, uses

RL: MOA (Modifier or additive use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(morphol. of acid-etched gel-spun polyethylene fibers)

L133 ANSWER 35 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1995:617867 Document No. 123:58736 Cardable hydrophobic
polyolefin fibers, process for their manufacture
and nonwoven textiles from. Thomsen, Lydia Dahl; Marcher,
Bjorn; Hansen, Pia Holm (Danaklon A/S, Den.). PCT Int. Appl. WO
9420664 A1 19940915, 48 pp. DESIGNATED STATES: W: AU, BR, CA, CN,
DE, DK, JP, KR, RU, UA, US; RW: AT, BE, CH, DE, DK, ES, FR, GB, GR,
IE, IT, LU, MC, NL, PT, SE. (English). CODEN: PIXXD2.
APPLICATION: WO 1994-DK91 19940304. PRIORITY: DK 1993-248 19930305.

AB A process for producing cardable, hydrophobic polyolefin
-based staple fibers, comprising applying to spun
filaments a first spin finish comprising
an antistatic agent, in particular a neutralized phosphoric acid
ester, stretching the filaments, applying to the stretched
filaments a second spin finish in the
form of a dispersion comprising an antistatic agent, a
natural or synthetic hydrocarbon wax with a m.p. 40-120°, or
a wax mixture comprising ≥1 hydrocarbon wax and having a m.p.

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40-120°, and optionally a polydiorganosiloxane,
     eg., di-Me siloxane, crimping the filaments,
     drying the filaments, and cutting the filaments
     to obtain staple fibers. The process allows the production of
     fibers that have controlled friction and hydrophobic
     properties and that can be carded at high speeds giving thermally
     bonded nonwovens having excellent hydrophobic
     characteristics. The nonwovens can function as liquid
     barriers in disposable diapers, feminine hygiene products, and
     medical gowns or drapes (no data).
     9002-88-4, Polyethylene
IT
     RL: PEP (Physical, engineering or chemical process); TEM (Technical
     or engineered material use); PROC (Process); USES (Uses)
        (fiber; cardable, hydrophobic and nonwoven
        textiles from)
     9002-88-4 HCAPLUS
RN
     Ethene, homopolymer (9CI) (CA INDEX NAME)
CN
     CM
     CRN
         74-85-1
     CMF C2 H4
H_2C = CH_2
IC
     ICM D06M013-292
     ICS D06M015-227
     40-7 (Textiles and Fibers)
CC
     Section cross-reference(s): 63
ST
     cardable hydrophobic polyolefin fiber finish;
    polypropylene cardable hydrophobic fiber finish;
    polyethylene cardable hydrophobic fiber finish; antistatic
     cardable hydrophobic polyolefin fiber; wax
     cardable hydrophobic polyolefin fiber;
     siloxane cardable hydrophobic polyolefin
     fiber
IT
    Antistatic agents
        (cardable hydrophobic polyolefin fibers,
        process for their manufacture and nonwoven textiles from)
IT
    Paraffin waxes and Hydrocarbon waxes, uses
    RL: NUU (Other use, unclassified); USES (Uses)
        (cardable hydrophobic polyolefin fibers,
        process for their manufacture and nonwoven textiles from)
IT
    Polypropene fibers, uses
    RL: PEP (Physical, engineering or chemical process); TEM (Technical
    or engineered material use); PROC (Process); USES (Uses)
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(cardable, hydrophobic and nonwoven textiles from)

## IT Polyolefin fibers

RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (process for manufacture of cardable, hydrophobic and nonwoven textiles from)

## IT Polyolefin fibers

RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (ethylene, cardable, hydrophobic and nonwoven textiles from)

IT 164715-25-7, Silastol F 203

RL: NUU (Other use, unclassified); USES (Uses)
(antistatic agent; for cardable hydrophobic polyolefin
fibers, process for their manufacture and nonwoven
textiles from)

IT 9002-88-4, Polyethylene 25085-53-4, Isotactic polypropylene

RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (fiber; cardable, hydrophobic and nonwoven textiles from)

IT 7664-38-2D, Phosphoric acid, C16-18 alkyl esters 9016-00-6, Dimethylsilanediol homopolymer, sru 31900-57-9, Dimethylsilanediol homopolymer

RL: NUU (Other use, unclassified); USES (Uses)
(for cardable hydrophobic polyolefin fibers,
process for their manufacture and nonwoven textiles from)

L133 ANSWER 36 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN

1995:220754 Document No. 122:216487 Polyether-containing fiber finishes. Doi, Tetsuo; Tatemoto, Shinichi; Kimura, Tsutomu; Munekyo, Takeshi (Matsumoto Yushi Seiyaku Kk, Japan). Jpn. Kokai Tokkyo Koho JP 06228885 A2 19940816 Heisei, 8 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1993-14605 19930201.

AB The fiber adhesion and fiber-fiber
abrasion resistance are improved by addition of (fatty acid-end-capped)
polyethers with mol. weight 1000-20,000 consisting of random
or block adducts of C4-24 alkylamines or C4-44 dialkylamines and
C2-4 alkylene oxides to finishes. Thus, a composition of isooctyl
palmitate 50, dioleyl adipate 10, polyoxyethylene oleyl ether 6,
ethoxylated hardened castor oil 10, oleic acid diethanolamide 5, Na
alkylsulfonate 5, KOAc 1, and laurylamine-ethylene oxide-propylene
oxide adduct (mol. weight 5000) 8 parts showed phase
inversion viscosity 1000 cP and its 10% aqueous
emulsion adhered strongly (16.8% as emulsion pickup) to
polyester filaments with good fiberfiber abrasion resistance (450 rubbing strokes until

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breakage).
IC
     ICM D06M015-53
CC
     40-7 (Textiles and Fibers)
    polyether fiber finish multifilament manuf; high speed
ST
     spinning fiber finish; alkoxylated
     alkylamine fiber finish; abrasion resistance polyether
     fiber finish; laurylamine polyoxyalkylene ether finish
    polyester fiber
IT
     Polyoxyalkylenes, uses
     RL: MOA (Modifier or additive use); USES (Uses)
        (amine ethers; fiber finishes containing (fatty
        acid-terminated) polyoxyalkylated aliphatic amines)
IT
     Polyester fibers, miscellaneous
     RL: MSC (Miscellaneous)
        (fiber finishes containing (fatty acid-terminated)
        polyoxyalkylated aliphatic amines)
IT
     Synthetic fibers, polymeric
    RL: MSC (Miscellaneous)
        (multifilaments; fiber finishes containing (fatty
        acid-terminated) polyoxyalkylated aliphatic amines)
IT
    Amines, uses
    RL: MOA (Modifier or additive use); USES (Uses)
        (aliphatic, alkoxylated; fiber finishes containing (fatty
        acid-terminated) polyoxyalkylated aliphatic amines)
IT
    Fatty acids, uses
    RL: MOA (Modifier or additive use); USES (Uses)
        (esters, with alkoxylated amines; fiber finishes containing
        (fatty acid-terminated) polyoxyalkylated aliphatic amines)
IT
    82497-16-3
                  84931-84-0
                              162031-77-8
                                             162031-99-4
                                                           162032-17-9
    162032-23-7, Ethylene oxide-propylene oxide
    copolymer ether with distearylamine
    RL: MOA (Modifier or additive use); USES (Uses)
        (fiber finishes containing (fatty acid-terminated)
       polyoxyalkylated aliphatic amines)
ΙT
    25038-59-9, Poly(ethylene terephthalate), miscellaneous
    RL: MSC (Miscellaneous)
        (filaments; fiber finishes containing (fatty
        acid-terminated) polyoxyalkylated aliphatic amines)
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L133 ANSWER 37 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1995:28115 Document No. 122:108475 Polybutylene monofilaments. I:
 Preparation and properties. Shaw, J. P.; Gilbert, M. (Inst. Polymer Technol. Mater. Eng., Loughborough Univ. Technol.,
 Loughborough/Leicestershire, LE11 3TU, UK). Plastics, Rubber and Composites Processing and Applications, 22(1), 9-27 (English) 1994.
 CODEN: PRPAEP. ISSN: 0959-8111.

AB The effect of processing variables, i.e., cooling water temperature,

AB The effect of processing variables, i.e., cooling water temperature, haul-off rate, and screw speed, on the properties of monofilaments

produced from 8 grades of 1-butene polymers was studied. crystallinity and birefringence of the monofilaments were measured, and wide-angle x-ray diffraction was used to indicate the extent of crystallite orientation. Tensile properties and elasticity were also measured. Except for low crystallinity grades and a highly nucleated variant, crystallinity increased with increasing water bath temperature, which resulted in increased modulus and elasticity. Break strain and break stress were reduced. For most samples, birefringence was almost independent of bath temperature, although increases with temperature were observed for three grades. Over a wide draw-ratio range, orientation behavior was shown to be complex and dependent on grade. Increased draw ratio generally increased break stress and modulus and decreased break strain. Stress at break was related to birefringence, although the nature of this relationship varied. Highly elastic materials showing up to 90% recovery were produced for certain grades by using appropriate processing conditions. High elasticity was associated with the combination of a highly developed crystalline phase, a high amorphous orientation, and a moderate crystalline orientation. 9002-88-4P, Polyethylene

IT

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

(butene-ethylene copolymer blends, fiber,

monofilament; effect of melt spinning conditions on properties. of)

RN9002-88-4 HCAPLUS

CN Ethene, homopolymer (9CI) (CA INDEX NAME)

> CM 1

CRN 74-85-1 CMF C2 H4

 $H_2C = CH_2$ 

CC 40-2 (Textiles and Fibers)

ST polybutene monofilament extrusion; tensile property polybutene fiber; crystallinity polybutene fiber; birefringence polybutene fiber; elasticity polybutene fiber; melt spinning butene polymer fiber

IT Polypropene fibers, preparation

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

(composite with butene copolymer, monofilament; effect of melt

spinning conditions on properties of) IT Polyolefin fibers RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (butene, monofilament; effect of melt spinning conditions on properties of) IT Polyolefin fibers RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (butene-ethylene, monofilament; effect of melt spinning conditions on properties of) IT Polyolefin fibers RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (ethylene, composite with butene copolymer, monofilament; effect of melt spinning conditions on properties of) IT 9002-88-4P, Polyethylene 9003-07-0P, Polypropylene RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (butene-ethylene copolymer blends, fiber, monofilament; effect of melt spinning conditions on properties of) IT 9003-28-5P, Poly(1-butene) RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (fiber, monofilament; effect of melt spinning conditions on properties of) L133 ANSWER 38 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN 1994:325703 Document No. 120:325703 Finish composition for fibers. Barsotti, Giampiero (Moplefan S.p.A., Italy). Eur. Pat. Appl. EP 576896 A2 19940105, 4 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE. (English). CODEN: EPXXDW. APPLICATION: EP 1993-109508 19930611. PRIORITY: IT 1992-MI1625 19920703. AB Title finish composition, especially for polyolefin fibers, comprises, sep. or as mixture, a fatty acid-polyalkyleneamine condensation product quaternized with a chlorohydrin-polyethylene glycol condensation product and a hydrogenated fatty acid-aminoalkylalkanolamine condensation product neutralized with an  $\alpha$ -hydroxycarboxylic acid, and containing an ethoxylated fatty alc. emulsifier. This composition gives good processability (antistatic effect) and high hydrophobicity to the fibers and good adhesion of adhesive labels to the nonwoven fabric articles produced from the treated fibers. IC ICM D06M013-46

ICS D06M013-402

40-7 (Textiles and Fibers)

CC

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ST
     polyolefin fiber spin finish
     ; polyalkylenamine fatty acid spin finish;
     propylene fiber spin finish;
     aminoalkylalkanolamine fatty acid spin finish;
     hydroxycarboxylic acid spin finish; alc fatty
     ethoxylated spin finish; antistatic
     polyolefin fiber spin finish
IT
     Polyolefin fibers
     Polypropene fibers, miscellaneous
     RL: USES (Uses)
        (finish composition for, with good processability and high
        hydrophobicity)
IT
    Alcohols, compounds
     RL: USES (Uses)
        (amino, reaction products, with fatty acids, finish composition, for
        polypropylene fibers, with good processability
        and high hydrophobicity)
IT
    Alcohols, uses
     RL: USES (Uses)
        (carboxy, finish composition, for polypropylene
        fibers, with good processability and high hydrophobicity)
IT
     Quaternary ammonium compounds, uses
    RL: USES (Uses)
        (fatty amido, finish composition, for polypropylene
        fibers, with good processability and high hydrophobicity)
IT
    Amides, compounds
    RL: USES (Uses)
        (fatty, quaternized, finish composition, for polypropylene
        fibers, with good processability and high hydrophobicity)
IT
     Tallow
    RL: USES (Uses)
        (hydrogenated, reaction products, with aminoethanolamine, finish
        composition, for polypropylene fibers, with good
        processability and high hydrophobicity)
IT
    Carboxylic acids, uses
    RL: USES (Uses)
        (hydroxy, finish composition, for polypropylene
        fibers, with good processability and high hydrophobicity)
IT
    Polyamines
    RL: USES (Uses)
        (polyalkylene-, reaction products, with fatty acids, quaternized,
        finish composition, for polypropylene fibers, with
        good processability and high hydrophobicity)
IT
    9003-07-0, Polypropylene
    RL: USES (Uses)
        (fiber, finish composition for, with good processability and
       high hydrophobicity)
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IT

57-11-4D, Stearic acid, reaction products with

tetraethylenepentamine 79-14-1, Glycolic acid, uses 109-84-2D, Aminoethanolamine, reaction products with hydrogenated tallow 112-57-2D, Tetraethylenepentamine, reaction products with stearic acid 25322-68-3D, Polyethylene glycol, fatty alkyl ethers 37569-89-4D, Polyethylene glycol bischlorohydrin ether, reaction products with fatty amides RL: USES (Uses)

(finish composition, for polypropylene fibers, with good processability and high hydrophobicity)

L133 ANSWER 39 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1994:220244 Document No. 120:220244 Analysis of ballistically caused damage in some test panel fibers. Cohen, S. H.; Prosser,
R. A.; King, A.; Desper, C. R. (Army Matick Res. Dev. Eng. Cent.,
MA, USA). Report, NATICK/TR-92/032; Order No. AD-A252220, 51 pp.
Avail. NTIS From: Gov. Rep. Announce. Index (U. S.) 1992, 92(20),
Abstr. No. 257,054 (English) 1992.

AB Microscopic and x-ray diffraction
methods to analyze ballistically induced damage to fibers
and yarns within multilayer panels of Spectra-900 and Spectra-1000,
polyethylene, and Kevlar are described. Microscopic evaluation by
SEM showed heat-induced damage in fibers several layers
beyond the layer in which the .22-caliber projectile came to rest,
and x-ray diffraction showed
crystallinity differences in fibers close to and
at some distance from the point of ballistic impact. Polarization
microscopy was used to corroborate the x-ray
diffraction findings, by elucidating differences in
birefringence within the fibers.

IT 9002-88-4, Polyethylene

RL: PRP (Properties)

(fibers, ballistically caused damage in panels of, testing of)

RN 9002-88-4 HCAPLUS

CN Ethene, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 74-85-1 CMF C2 H4

 $H_2C = CH_2$ 

CC 40-4 (Textiles and Fibers)
ST ballistic damage test panel

ST ballistic damage test panel fiber; polyethylene fiber ballistic damage panel; polyester fiber

ballistic damage panel; aramid fiber ballistic damage panel

IT Testing of materials

(ballistic, of fiber panels)

IT Polyester fibers, properties

RL: PRP (Properties)

(ballistically caused damage in panels of, testing of)

IT Polyamide fibers, properties

RL: PRP (Properties)

(aramid, ballistically caused damage in panels of Kevlar, testing of)

IT Polyolefin fibers

RL: PRP (Properties)

(ethylene, ballistically caused damage in panels of Spectra 900, testing of)

IT 9002-88-4, Polyethylene

RL: PRP (Properties)

(fibers, ballistically caused damage in panels of, testing of)

L133 ANSWER 40 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN

- 1994:193943 Document No. 120:193943 Process for high-speed spinning of polyester fibers with reduced neps and yarn breakage.

  Makino, Shoji; Taniguchi, Katsutoshi (Teijin Ltd., Japan). PCT Int. Appl. WO 9320268 A1 19931014, 21 pp. DESIGNATED STATES: W: KR, US; RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE. (Japanese). CODEN: PIXXD2. APPLICATION: WO 1993-JP398 19930330. PRIORITY: JP 1992-79771 19920401.
- AB In the title process, spun polyester fibers are coated with aqueous lubricant emulsions containing mainly monobasic acid esters having average mol. weight 300-500, 1-15% polyoxyalkylene glycol copolymers having average mol. weight ≥1000, and 0.1-3% organic siloxane compds. and/or fluoroalkyl group-containing compds. and wound at ≥3000 m/min. PET was melt spun, coated with 10% emulsion containing octyl stearate 60, ethylene oxide-polytetramethylene glycol copolymer 5, and ethylene oxide-modified silicone 2 parts to effective component content 0.4%, taken up at 4000 m/min, and drawn to give fibers with low number of neps and yarn breaks.

IC ICM D01F006-62

ICS D01F011-08; D06M015-53; D06M013-224; D06M013-50; D06M013-08

ICI D06M101-32

CC 40-2 (Textiles and Fibers)

ST polyester fiber high speed spinning; octyl stearate lubricant finish polyester spinning; polyoxyalkylene lubricant finish polyester spinning; silicone lubricant finish polyester spinning; lubricant finish polyester high speed spinning

IT Lubricants

(finishes, containing monobasic acid esters, polyoxyalkylenes and silicones and/or fluoroalkyl group-containing compds., for high-speed spinning of polyester fibers with reduced neps)

IT Polyoxyalkylenes, uses

RL: TEM (Technical or engineered material use); USES (Uses) (lubricant finishes containing, for high-speed spinning of polyester fibers with reduced neps)

IT Polyester fibers, preparation

RL: SPN (Synthetic preparation); PREP (Preparation) (spinning of, at high speeds, lubricant finishes containing monobasic acid esters, polyoxyalkylenes and silicones and/or fluoroalkyl group-containing compds. for)

IT Siloxanes and Silicones, uses

RL: TEM (Technical or engineered material use); USES (Uses)
(amino-containing, lubricant finishes containing, for high-speed spinning of polyester fibers with reduced neps)

IT Siloxanes and Silicones, compounds

RL: TEM (Technical or engineered material use); USES (Uses)
(ethoxylated, monobasic acid ester lubricant finishes containing, for high-speed spinning of polyester fibers with reduced neps)

IT Acids, compounds

RL: TEM (Technical or engineered material use); USES (Uses) (monobasic, esters, lubricant finishes containing, for high-speed spinning of polyester fibers)

IT Ethers, uses

RL: TEM (Technical or engineered material use); USES (Uses) (perfluoroalkyl, monobasic acid ester lubricant finishes containing, for high-speed spinning of polyester **fibers** with reduced neps)

IT Siloxanes and Silicones, uses

RL: TEM (Technical or engineered material use); USES (Uses) (polyester-, lubricant finishes containing, for high-speed spinning of polyester fibers with reduced neps)

IT Siloxanes and Silicones, uses

RL: TEM (Technical or engineered material use); USES (Uses) (polyether-, lubricant finishes containing, for high-speed spinning of polyester fibers with reduced neps)

IT Polyesters, uses

RL: TEM (Technical or engineered material use); USES (Uses) (siloxane-, lubricant finishes containing, for high-speed spinning of polyester fibers with reduced neps)

IT 25038-59-9P, Poly(ethylene terephthalate), preparation RL: SPN (Synthetic preparation); PREP (Preparation)

(fiber, high-speed spinning of, with reduced neps and yarn breakage, lubricant finishes for)

IT 109-36-4, Octyl stearate 2306-92-5, Octyl decanoate 3687-45-4,

```
13945-76-1, Lauryl laurate 16958-85-3, Octyl
Oleyl oleate
palmitate 22047-49-0, Octyl stearate 70364-64-6, Isotridecyl
           72576-78-4
palmitate
RL: TEM (Technical or engineered material use); USES (Uses)
   (lubricant finishes containing, for high-speed spinning of polyester
   fibers with reduced neps)
9003-11-6, Ethylene oxide-propylene oxide
copolymer
           25322-68-3D, Polyethylene glycol, ethers with
           27637-03-2
silicones
RL: TEM (Technical or engineered material use); USES (Uses)
   (monobasic acid ester lubricant finishes containing, for high-speed
   spinning of polyester fibers with reduced neps)
```

L133 ANSWER 41 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN 1993:651245 Document No. 119:251245 Polyethylene fibers -polyethylene matrix composites: preparation and physical properties. Teishev, Albert; Incardona, Silvia; Migliaresi, Claudio; Marom, Gad (Grad. Sch. Appl. Sci. Technol., Hebrew Univ., Jerusalem, 91904, Israel). Journal of Applied Polymer Science, 50(3), 503-12 (English) 1993. CODEN: JAPNAB. ISSN: 0021-8995. Drawing on the difference in m.ps. of ultrahigh-modulus AΒ polyethylene (I) fiber (150°) and HDPE matrix (130°), single-polymer composites were fabricated under various processing conditions. Because of the chemical similarity of the composite components, good bonding at the fiber-matrix interface could be expected. The matrix, the fiber, and unidirectional composite laminate were studied using thermomech. and DSC analyses, a hot-stage crystallization unit attached to a polarizing microscope, and a universal tensile testing machine. thermomech. anal. showed neg. thermal expansion of the fiber over the complete temperature range of the experiment Three regimes of contraction according to the values of the thermal expansion coefficient were detected. DSC analyses of either the fiber or the composite specimens did not show any appreciable changes after various thermal treatments. They also showed no evidence of fiber relaxation during manufacture, probably because of the pressure-related transverse constraint. The tensile strength and modulus values of the composite appeared to be fairly high and close to those reported for other composites reinforced with I fibers. A study of the matrix microstructure did not give any proof of transcryst. growth at the fiber-matrix interface even for chemical or plasma surface-treated fibers.

IT 9002-88-4P, Polyethylene
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (fiber-matrix composites, preparation and properties of)
RN 9002-88-4 HCAPLUS

CN Ethene, homopolymer (9CI) (CA INDEX NAME)

IT

CM 1

CRN 74-85-1 CMF C2 H4

 $H_2C = CH_2$ 

CC 37-5 (Plastics Manufacture and Processing)

ST polyethylene **fiber** matrix composite; expansion contraction polyethylene composite; strength polyethylene **fiber** matrix composite

IT Crystallinity

(of polyethylene fiber-matrix composites, properties in relation to)

IT Polyolefin fibers

RL: SPN (Synthetic preparation); PREP (Preparation) (ethylene, HDPE composites, preparation and properties of)

IT 9002-88-4P, Polyethylene

RL: SPN (Synthetic preparation); PREP (Preparation) (fiber-matrix composites, preparation and properties of)

L133 ANSWER 42 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1993:605418 Document No. 119:205418 Finish for textile fibers
. Petrea, Randy Dell; Schuette, Robert Louis (Milliken Research Corp., USA). Eur. Pat. Appl. EP 547846 A1 19930623, 11 pp.
DESIGNATED STATES: R: BE, DE, FR, GB, IT, NL. (English). CODEN:
EPXXDW. APPLICATION: EP 1992-311346 19921211. PRIORITY: US
1991-806990 19911213.

AB Title finish comprises poly- $\alpha$  -olefins 70-95, an emulsifier selected from ethoxylated C12-36 branched alcs., alkoxylated ≥3 OH group containing polyhydric alcs. which are esterified with C12-36 fatty acids, and alkoxylated glycerol esters of C12-36 fatty acids having ≥1 OH group 5-30, ionic emulsifiers ≤10, and antisoiling additive ≤5 parts. These compns. are used as spin finishes for synthetic fibers, especially elastomeric polyurethanes, polyesters, and polyamides. A composition containing of a 4 cSt poly(. alpha.-olefin) 80, ethoxylated 2-octyldodecanol (7 EO) 20, C12-15 5 ethylene oxide phosphate 5.3, and ethoxylated castor oil 4.5 g was prepared and 7.9 g H2O added to provide a clear stable mixture This composition on 70/34 polyester had fiber/ fiber hydrodynamic friction 0.57 and boundary kinetic fiber/fiber friction 0.14.

IC ICM D06M013-03

ICS D06M015-227; D06M013-17; D06M013-165; D06M013-224

CC 40-7 (Textiles and Fibers)

```
ST
     spin finish polyolefin synthetic
     fiber; alkoxylated alc spin finish;
     polyhydric alc alkoxylated spin finish; glycerol
     ester spin finish
     Emulsifying agents
IT
        (in manufacture of spin finishes for synthetic
        fibers)
IT
     Polyamide fibers, miscellaneous
     Polyester fibers, miscellaneous
     Spandex fibers
     RL: MSC (Miscellaneous)
        (spin finishes for)
IT
     Fatty acids, esters
     RL: USES (Uses)
        (C12-36, esters, with polyhydric alcs., ethoxylated, in manufacture of
        spin finishes for synthetic fibers)
IT
     Alcohols, compounds
     RL: USES (Uses)
        (C12-36, ethoxylated, in manufacture of spin
        finishes for synthetic fibers)
İT
     Castor oil
     Glycerides, compounds
     RL: USES (Uses)
        (ethoxylated, in manufacture of spin finishes for
        synthetic fibers)
     Alkenes, polymers
IT
     RL: USES (Uses)
        (\alpha-, polymers, oligomers, spin finish
        compns. for synthetic fibers containing)
     9003-27-4, Tebeflex 200
IT
                               9056-42-2D, C12-15-alkyl ethers
                  150825-14-2
     32128-65-7
     RL: USES (Uses)
        (in manufacture of spin finishes for synthetic
        fibers)
ĬΤ
     872-05-9, 1-Decene
                          1120-36-1, 1-Tetradecene
                                                      25067-08-7,
     1-Dodecene homopolymer 25068-25-1, 1-Octene homopolymer
     RL: USES (Uses)
        (oligomers, in manufacture of spin finishes for
        synthetic fibers)
L133 ANSWER 43 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1990:22298
             Document No. 112:22298 Spinning finish
     for polypropylene and polyamide fibers for
     improved processability. Svoboda, Karol; Mikolaj, Dusan; Babjak,
     Jan; Simo, Rudolf; Nemeth, Ladislav (Czech.). Czech. CS 260758 B1
     19890515, 5 pp.
                      (Slovak). CODEN: CZXXA9. APPLICATION: CS
     1985-5353 19850719.
```

An 1-25% aqueous dispersion of tech. olefin

AB

ethoxylated with 10-30 mol oxirane is applied to polypropylene or polyamide fibers before the spinneret as a finish for effectively reducing the friction between fibers and metal or corundum and improving coherence of a tow during winding, stretching, texturing, and other operations. The content of finish on fibers is 0.3-2.2%.

IC ICM D06M015-263

CC 40-7 (Textiles and Fibers)

ST olefin ethoxylated finish polypropylene fiber; polyamide fiber finish ethoxylated olefin; friction redn polypropylene fiber; processability mech polypropane fiber

IT Lubricants

(ethoxylated olein, for finishing of polyamide or polypropene fibers for improved processability)

IT Polyamide fibers, uses and miscellaneous Polypropene fibers, uses and miscellaneous RL: USES (Uses)

(spinning finishes for, ethoxylated olein as, for reduced friction and improved processability)

IT 68958-64-5

RL: USES (Uses)

(finishes, for polyamide or polypropylene fibers, for reduced friction)

L133 ANSWER 44 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1987:139494 Document No. 106:139494 Polyester molding compositions.
Okamoto, Takashi; Ohama, Fumio (Unitika Ltd., Japan). Jpn. Kokai
Tokkyo Koho JP 61241353 A2 19861027 Showa, 8 pp. (Japanese).
CODEN: JKXXAF. APPLICATION: JP 1985-85167 19850418.

AB The title compns., having low-temperature impact strength, heat resistance, and good processability, comprise a poly(ethylene terephthalate) (I)-polyarylate mixture 100, an inorg. nucleating agent having average particle diameter ≤50 µ and/or metal carboxylate-containing organic compound (or polymer) 0.05-10, a polyolefin or elastomer modified with 0.001-10 mol % alicyclic carboxylic acid or its derivative 3-30,  $80-99:1-20:0-19 \alpha$  -olefin-glycidyl (meth)acrylate-vinyl acetate copolymer 1-30, and reinforcing fibers 0-150 parts. Thus, a mixture of 70% I (intrinsic viscosity 0.68; 6:4 PhOH-C2H2Cl4; 0.5%; 20°) and 30% polyarylate (prepared from 1:1 terephthalic acid-isophthalic acid mixture and bisphenol A; reduced viscosity 0.65; 25°; 1 g/dL) 100, Surlyn 1555 (II) 3, modified copolymer (III) (prepared from 72.0:28.0 ethylene-propene rubber 1000, endo-bicyclo[2.2.1]hept-5ene-2,3-dicarboxylic anhydride 3, and tert-Bu2O2 1 part) 10, and 92:8 ethylene-glycidyl methacrylate copolymer 5 parts was molded at

260° with molding cycle 20 s to give a molding having Izod impact strength (kg-cm/cm) 38 at room temperature and 23 at -30° and heat-distortion temperature 85°, vs. 15, 6, and 70, resp., without II and III. 9002-88-4D, Polyethylene, carboxylated IT RL: USES (Uses) (aromatic polyester blends containing, heat- and impact-resistant) 9002-88-4 HCAPLUS RN Ethene, homopolymer (9CI) (CA INDEX NAME) CN CM CRN 74-85-1 CMF C2 H4  $H_2C = CH_2$ IT 14807-96-6, Talc, uses and miscellaneous RL: USES (Uses) (nucleating agents, aromatic polyester blends containing) 14807-96-6 HCAPLUS RN CN Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME) HO-Si-OH ●3/4 Mg IC ICM C08L067-02 ICS C08K003-00; C08K005-09; C08K013-04 C08L067-02, C08L023-26, C08L023-02; C08L067-02, C08L023-26, C08L063-00; C08K013-04, C08K007-14, C08K003-00; C08K013-04, C08K007-14, C08K005-09 38-3 (Plastics Fabrication and Uses) CC ST polyarylate blend polyethylene terephthalate; polyester blend molding; nucleating agent polyester blend; ionomer polyester blend; bicycloheptenedicarboxylic polyolefin polyester blend; ethylene copolymer polyester

blend; glycidyl acrylate copolymer blend; vinyl acetate copolymer blend; impact strength polyester blend; heat resistance polyester

blend; bisphenol benzenedicarboxylic polyester blend IT Kaolin, uses and miscellaneous Mica-group minerals, uses and miscellaneous RL: USES (Uses) (nucleating agents, aromatic polyester blends containing) IT Crystal nucleation (agents, carboxylate compds., aromatic polyester blends containing) 129-64-6D, reaction products with olefin polymers IT 9002-88-4D, Polyethylene, carboxylated 9003-07-0D, Polypropylene, carboxylated 24937-78-8D, Ethylene-vinyl acetate copolymer, carboxylated 25038-59-9, Poly(ethylene terephthalate), uses and miscellaneous 25087-34-7D, 1-Butene-ethylene copolymer, carboxylated 26061-90-5, Ethylene-glycidyl methacrylate copolymer RL: USES (Uses) (aromatic polyester blends containing, heat- and impact-resistant) IT 7631-86-9, Silica, uses and miscellaneous 14807-96-6, Talc, uses and miscellaneous RL: USES (Uses) (nucleating agents, aromatic polyester blends containing) L133 ANSWER 45 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN 1986:462170 Document No. 105:62170 Blends of fluoroalkylguanidines with fluorinated polyoxyalkylenes. Chang, John C.; Williams, Kathryn L. (Minnesota Mining and Manufacturing Co., USA). U.S. US 4565641 A 19860121, 13 pp. Cont.-in-part of U.S. Ser. No. 440,317, abandoned. (English). CODEN: USXXAM. APPLICATION: US 1984-633977 19840724. PRIORITY: US 1982-440317 19821109. · AB Mixts. of (fluoroalkyl) guanidines and polyoxyalkylenes bearing fluoroalkyl groups are useful in organic solvents or as aqueous dispersions in oil- and waterproofing fibers. Thus, a neat oil-spin finish containing 13.1% RZ[NHC(NBu2):NZ]2R[R = C8F17SO2N(Et)CH2CH2OCONH, Z = CH2(C6H4)2],5.0% acrylic polymer [from C8F17SO2N(Bu)CH2CH2OCOCH:CH2 and polyethylene-polypropylene glycol mono- and diacrylate] 46.2% coconut oil-based lubricant, and 35.7% Bu(OCH2CH2)2OH was coated (425 ppm F) on nylon fibers which were spun to a carpet retaining 81% F after dyeing and having good resistance to oil, water, and walk-on soiling. IC ICM D06M013-08 INCL 252008750 40-9 (Textiles) CC

fluoropolymer finish fiber; waterproofing finish

fiber; soilproofing finish fiber; oilproofing
finish fiber; guanidine fluoroalkyl finish fiber

ST

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; acrylate copolymer finish fiber; polyoxyalkylene
     fluoropolymer finish fiber; nylon carpet finish
IT
     Carpets
     Polyamide fibers, uses and miscellaneous
     RL: USES (Uses)
        (water- and oilproofing agents for, (fluoroalkyl)guanidines and
        fluorinated polyoxyalkylenes as)
L133 ANSWER 46 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
             Document No. 103:88990 Spin finish
1985:488990
     compositions. Anon. (UK). Research Disclosure, 253, 250 (No.
     25342) (English) 1985. RD 253042 19850510. CODEN: RSDSBB. ISSN:
     0374-4353. PRIORITY: RD 1985-253042 19850510.
AB
     The addition of 0.5-5 weight% of alkyl glycidyl esters of polycarboxylic
     acid or anhydride-containing adducts of olefinically unsatd.
     natural oils and of polycarboxylic acid- or anhydride-containing
     copolymers with \alpha -olefin compds.
     to spin finish compns. for polyamide or .
     polyester yarns comprising a glycerol triester, a sulfated glycerol
     triester, a salt of a partial phosphate ester, and a salt of a
     higher fatty acid is disclosed. Alternatively, use may be made of
     the alkyl alkoxyalkyl esters and mixed esters of copolymers of
     maleic anhydride with C2-30 (preferably C12-24) \alpha -
     olefins.
CC
     40-7 (Textiles)
ST
     spin finish polyamide fiber; polyester
     fiber spin finish; polycarboxylic acid
     spin finish fiber; olefin
     copolymer spin finish fiber;
     maleic anhydride polymer spin finish; glycidyl
     ester spin finish fiber
IT
     Polyamide fibers, uses and miscellaneous
     Polyester fibers, uses and miscellaneous
     RL: USES (Uses)
        (spin finish compns. for)
IT
     Alkenes, polymers
     RL: USES (Uses)
        (C12-24, polymers with maleic anhydride, esters, spin
        finish compns., for polyamide and polyester
        fibers)
IT
     Carboxylic acids, compounds
     RL: USES (Uses)
        (poly-, adducts with natural oils, alkyl glycidyl esters,
        spin finish compns., for polyamide and
        polyester fibers)
IT
    108-31-6D, polymers with \alpha -olefins
     , esters
    RL: USES (Uses)
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(spin finish compns., for polyamide and polyester fibers)

L133 ANSWER 47 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN 1985:221679 Document No. 102:221679 Fire-resistant polyester molding compositions. (Unitika Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 60015451 A2 19850126 Showa, 13 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1983-123597 19830706. AB A fire-resistant, glass fiber-reinforced polyester composition, giving a dimensionally stable product at a fast molding cycle, comprises a polyester with ≥80% ethylene terephthalate repeating units 100, a nucleating agent 0.1-5, a nucleating aid 0.5-10, and an organic halogenated compound 10-30 parts, in addition to an Sb-containing compound to give 0.2-1:1 Sb compound-halogen compound ratio, a glass fiber at 10-15 parts/composition, and an inorg. filler at 15-45 parts/composition Thus, a pelletized blend comprising poly(ethylene terephthalate) [25038-59-9] 10, Himilan 1555 (ethylene-methacrylic acid copolymer Na salt) [25608-26-8] nucleating agent 4, 72:28 ethylenepropylene copolymer modified with endo-bicyclo-[2,2,1]-5-heptene-2,3-dicarboxylic anhydride (nucleating aid) 5, GLC 935 P [poly(dibromophenylene oxide)] [74082-93-2] 16, and Sb2O3 5 parts, in addition to 20% powdered talc and 10% glass fiber (average fiber length 0.325 mm), was injected at 600-700 kg/cm2 and 240-250° into a mold at 100-105° at molding time 20 s to give a sample exhibiting crystallization temperature 116.2°, heat-distortion temperature 207°, flexural strength 1250 kg/cm2, flexural modulus 66,000 kg/cm2, warpage (sample thickness 1.6 mm, after heat treatment at 130° for 2 h) 5.26-5.50 mm, and flame time (UL 94) <10 s, with no blooming of fire retardants. IT 14807-96-6, uses and miscellaneous RL: USES (Uses) (fillers, for glass fiber-reinforced polyesters, with

RN CN low molding warpage)

Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)

14807-96-6 HCAPLUS

●3/4 Mg

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IC
     ICM C08L067-02
     ICS
         C08K013-04
     C08K013-04, C08K007-14, C08K003-00, C08K005-00
ICI
     37-6 (Plastics Manufacture and Processing)
CC
ST
     polyethylene terephthalate glass fiber reinforcement;
     ionomer nucleating agent polyester molding;
     bicycloheptenedicarboxylic anhydride graft polyolefin;
     polydibromophenylene oxide fireproofing agent polyester;
     talc filler polyester molding
     Crystal nucleation
IT
        (agents, ionomers and carboxy-grafted
        polyolefins, for glass fiber-polyester
        moldings)
IT
     Polyoxyphenylenes
     RL: USES (Uses)
        (brominated, fireproofing agents, for glass fiber
        -reinforced polyesters, with low blooming)
IT
     Ionomers
     RL: USES (Uses)
        (ethylene-methacrylic acid copolymer sodium salts,
        nucleating agents, for glass fiber
        -reinforced polyesters moldable at fast molding cycles, with low
        product warpage)
    Mica-group minerals, uses and miscellaneous
IT
     RL: USES (Uses)
        (fillers, for glass fiber-reinforced polyesters, with
        low molding warpage)
IT
     Polyesters, uses and miscellaneous
     RL: USES (Uses)
        (glass fiber-reinforced, fire-resistant, moldable at
        fast molding cycles, with low product warpage)
IT
     Fireproofing agents
        (poly(dibromophenylene oxide), for glass fiber
        -reinforced polyester moldings, with low blooming)
IT
     1344-95-2 14807-96-6, uses and miscellaneous
     RL: USES (Uses)
        (fillers, for glass fiber-reinforced polyesters, with
        low molding warpage)
IT
    74082-93-2
    RL: USES (Uses)
        (fireproofing agents, for glass fiber-reinforced
        polyester moldings, with low blooming)
IT
     9016-88-0
                 25038-59-9, uses and miscellaneous
    RL: USES (Uses)
        (glass fiber-reinforced, fire-resistant, moldable at
        fast molding cycles, with low product warpage)
```

IT

25608-26-8

(nucleating agents, for glass fiber -reinforced polyesters, moldable at fast molding cycles, with low product warpage) L133 ANSWER 48 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN 1984:211031 Document No. 100:211031 Polyester molding compositions. (Mitsubishi Chemical Industries Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 58187449 A2 19831101 Showa, 5 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1982-70657 19820427. Polyester molding compns. containing inorg. reinforcing fillers 5-150, AB crystallization accelerators (C15-400 fatty acid salts, silicates, or (meth)acrylic acid-olefin copolymer Na or K salts) 0.1-10, triaryl phosphates 0.5-10, and tris(2,4-di-tertbutylphenyl) phosphite (I) [31570-04-4] or trilauryl trithiophosphite [1656-63-9] 0.001-10 phr have good thermal stability and crystallinity. Thus, 185 g EtCO2H was alkylated with 167.5 g C≥30 1-alkene mixture (peroxide catalyzed) to give a fatty acid which was converted to its Na salt (II). An injection molding composition was prepared from poly(ethylene terephthalate) [25038-59-9] 66.5, glass fibers 30, II 1.5, tricresyl phosphate [1330-78-5] 1.5, and I 0.5 part. IT 14807-96-6, uses and miscellaneous RL: USES (Uses) (nucleating agents, for polyester molding compns.) RN 14807-96-6 HCAPLUS CN Talc (Mq3H2(SiO3)4) (9CI) (CA INDEX NAME)

## ●3/4 Mq

IC

CC 37-6 (Plastics Manufacture and Processing)
ST polyester molding compn; phosphite ester heat stabilizer; phosphate
 ester heat stabilizer; heat stabilizer polyester molding;
 nucleating agent polyester molding
IT Crystal nucleation

(agents, for polyester molding compns.)

C08L067-02; C08K005-51; C08K007-02; C08K007-16

IT Ionomers
RL: USES (Uses)

RL: USES (Uses)

```
(nucleating agents, for polyester molding
        compns.)
IT
     Carboxylic acids, compounds
     RL: USES (Uses)
        (C15-400, salts, nucleating agents, for
        polyester molding compns.)
IT
     25038-59-9, uses and miscellaneous
     RL: USES (Uses)
        (molding compns., nucleating agents and heat
        stabilizers for)
İT
     79-09-4D, alkyl derivs., sodium salts 14807-96-6, uses and
     miscellaneous
                     25608-26-8
     RL: USES (Uses)
        (nucleating agents, for polyester molding
L133 ANSWER 49 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
             Document No. 100:211030 Polyester molding compositions.
1984:211030
     (Mitsubishi Chemical Industries Co., Ltd., Japan). Jpn. Kokai
     Tokkyo Koho JP 58187448 A2 19831101 Showa, 7 pp. (Japanese).
     CODEN: JKXXAF. APPLICATION: JP 1982-70656 19820427.
     The title compns., containing polyolefins 0.5-50, crystallization
AB
     accelerators (C15-400 carboxylic acid salts, silicates, or
     (meth)acrylic acid-olefin copolymer salts)
     0.1-10, triaryl phosphates 0.5-10, and tris(2,4-di-tert-butylphenyl)
     phosphite (I) [31570-04-4], trilauryl trithiophosphite
     [1656-63-9], or 2,2-thiodiethylene bis[3-(3,5-di-tert-4-
     hydroxyphenyl)propionate] [41484-35-9] 0.001-10 phr have good
     impact resistance, heat resistance, and crystallinity. Thus, 185 g
     EtCO2H was alkylated with 167.5 parts Dialene 30 (C>30 1-alkene
    mixture) in the presence of a peroxide to give a fatty acid, which was
     converted to its Na salt (II). An injection molding composition was
    prepared from poly(ethylene terephthalate) [25038-59-9] 61.5, glass
     fibers 30, polyethylene [9002-88-4] 5, II 1.5,
     tricresyl phosphate [1330-78-5] 1.5, and I 0.5 part.
IT
     9002-88-4 14807-96-6, uses and miscellaneous
    RL: USES (Uses)
        (nucleating agent, for polyester molding
       compns.)
RN
     9002-88-4 HCAPLUS
CN
    Ethene, homopolymer (9CI) (CA INDEX NAME)
    CM
         1
     CRN 74-85-1
    CMF C2 H4
```

 $H_2C = CH_2$ RN 14807-96-6 HCAPLUS CN Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME) 0 HO-Si-OH ●3/4 Mg IC C08L067-02; C08K005-51 C08L067-02, C08L023-08, C08L033-02, C08L051-06 ICI CC 37-6 (Plastics Manufacture and Processing) ST polyester compn injection molding; nucleating agent polyester molding; heat stabilizer polyster molding; phosphate heat stabilizer; phosphite heat stabilizer; polyethylene nucleating agent Crystal nucleation (agents, for polyester molding compns.) IT Ionomers RL: USES (Uses) (nucleating agent, for polyester molding compns.) IT Carboxylic acids, compounds RL: USES (Uses) (C15-400, salts, nucleating agents for polyester molding compns.) IT 25038-59-9, uses and miscellaneous RL: USES (Uses) (molding compns., nucleating agents and stabilizers for) IT 79-09-4D, alkyl derivs., sodium salts 9002-88-4 14807-96-6, uses and miscellaneous 25608-26-8 36704-47-9 RL: USES (Uses) (nucleating agent, for polyester molding

L133 ANSWER 50 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1984:105053 Document No. 100:105053 Durable antisoiling coatings for textile filaments. Champaneria, Nitin J.; Harper, Lee R.; Hosegood, Edward A. (du Pont de Nemours, E. I., and Co. , USA).
U.S. US 4388372 A 19830614, 10 pp. Cont.-in-part of U.S. 4,325,857.

compns.)

(English). CODEN: USXXAM. APPLICATION: US 1982-356716 19820310. PRIORITY: US 1980-149334 19800513; BR 1981-2732 19810505. AB Durable antisoiling compns. for textiles contain perfluoroalkyl esters of citric acid urethanes and reaction products of carboxylated vinyl polymers, epoxy resins, and tertiary amines. compns. can be applied as aqueous dispersions containing nonionic textile lubricants. Thus, nylon 66 carpet yarn was finished with a mixture of polyethylene glycol lubricant containing 4.5% diurethane from F(CF2)6-14CH2CH2OH triester with citric acid and hexamethylene diisocyanate, and a condensate of Me2NCH2CH2OH, epichlorohydrin-bisphenol A polymer, and Et acrylate-methacrylic acid-styrene polymer. A carpet from this yarn has a soil rating of 4, compared with 6 for an untreated carpet and 3 for a topically treated carpet. A primary spin finish for the manufacture of bulk continuous-filament nylon 66 carpet yarn in a coupled spin-draw-bulk process is prepared by adding 165.4 lbs. of a 99% concentrate of ethylene oxidepropylene oxide copolymer [9003-11-6] to 1310 lbs. water followed by slow addition of 39.4 lbs. 50% aqueous dispersion of ethoxylated castor oil, slowly adding with mixing 281 lbs. of a 5.6% dispersion of an acrylic-modified resin, adding 184.9 lbs. of a 51% aqueous dispersion of a fluoro ester citric acid urethane, and diluting with water to give a composition with solid content 14.25%. The polyamide yarn was treated with 0.45% finish, drawn, bulked, treated with a secondary finish based on coconut oil and ethoxylated castor oil, tufted into a carpet backing, and dyed to give a fabric very resistant to soiling under normal foot traffic. B32B027-34; B32B027-36; C08L063-00

IC

INCL 428395000

40-7 (Textiles) CC

IT Epoxy resins, uses and miscellaneous RL: USES (Uses)

> (aminated, reaction products with carboxylated polymers, in antisoiling compns. for polyamide fibers)

IT Polyamide fibers, uses and miscellaneous

RL: USES (Uses)

(antisoiling compns. for)

ΙT Lubricants

> (for polyamide fibers, containing durable antisoiling finishes)

IT Soilproofing

(of polyamide fibers, lubricant finishes containing)

IT Polyamide fibers, uses and miscellaneous RL: USES (Uses)

(soilproofing composition for carpet yarn from)

IT 77-92-9D, fluoroalkyl esters, urethanes 108-01-0D, reaction products with epoxy resins and methacrylic acid copolymers

25035-68-1D,

25068-38-6D, reaction

822-06-0D, urethanes with tris(fluoroalkyl) citrates

reaction products with aminated epoxy resins

products with amines and methacrylic acid copolymers RL: USES (Uses) (soilproofing finishes containing, for nylon fibers) L133 ANSWER 51 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN Document No. 98:180690 Poly(ethylene terephthalate) 1983:180690 molding compositions. (Dainippon Ink and Chemicals, Inc., Japan). Jpn. Kokai Tokkyo Koho JP 57187350 A2 19821118 Showa, 5 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1981-72246 19810515. AB Molding compns. comprise poly(ethylene terephthalate) (I) [25038-59-9] 100, olefin copolymers containing carboxylic acid metal salts (for nucleating agents) 0.1-15, aromatic polyester polyether elastomers (for lowering crystallization beginning temps.) 0.5-25, and polyfunctional compds. selected from polycarboxylic acid anhydrides, polyepoxides, and polyisocyanates 0-10 parts. Thus, test pieces prepared from I 100, Himilan 1601 (ethylene-methacrylic acid copolymer Na salt) [84683-11-4] 2, Pelprene P70B 10, glass fibers 20, and wollastonite 80 parts had good surface gloss, heat distortion temperature 201°, and crystallization beginning temperature 100°, compared with very poor, 64, and 119, resp., for test pieces prepared from I alone. 14807-96-6, uses and miscellaneous RL: USES (Uses) (fillers, containing glass fibers, for poly(ethylene terephthalate) containing ethylene-methacrylic acid copolymer sodium salt and polyester-polyether elastomers) 14807-96-6 HCAPLUS RN CN Talc (Mg3H2(SiO3)4) (9CI) (CA INDEX NAME)

HO-Si-OH

## ●3/4 Mg

IC C08L067-02; C08L033-02
ICA C08K005-10; C08K005-29; C08L063-00
CC 38-3 (Plastics Fabrication and Uses)
ST polyethylene terephthalate nucleating agent;
 polyether rubber polyester molding; glass fiber polyester
 molding; wollastonite polyester molding

IT Crystal nucleation (agents, ethylene-methacrylic acid copolymer sodium salts, for poly(ethylene terephthalate) containing polyester-polyether elastomers) Ionomers IT RL: USES (Uses) (nucleating agents, for poly(ethylene terephthalate) containing polyester-polyether elastomers) IT Rubber, synthetic RL: USES (Uses) (polyester-polyether, poly(ethylene terephthalate) containing ethylene-methacrylic acid copolymer sodium salt and glass fibers and wollastonite and, for moldings) IT 13983-17-0 14807-96-6, uses and miscellaneous RL: USES (Uses) (fillers, containing glass fibers, for poly(ethylene terephthalate) containing ethylene-methacrylic acid copolymer sodium salt and polyester-polyether elastomers) IT 25038-59-9, uses and miscellaneous RL: USES (Uses) (moldings, containing ethylene-methacrylic acid copolymer sodium salt and polyester-polyether elastomers and glass fibers and wollastonite) IT 25608-26-8 RL: USES (Uses) (nucleating agents, for poly(ethylene terephthalate) containing polyester-polyether elastomers) L133 ANSWER 52 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN Document No. 97:146183 Lubricant compositions for 1982:546183 finishing synthetic fibers. Decker, Quintin William; Marcus, Erich (Union Carbide Corp., USA). Eur. Pat. Appl. EP 54953 Al 19820630, 23 pp. DESIGNATED STATES: R: BE, DE, FR, GB, IT, NL, (English). CODEN: EPXXDW. APPLICATION: EP 1981-110657 PRIORITY: US 1980-219217 19801222. 19811221. AB Spin finishes for synthetic fibers having good thermal stability and low fume emission contain 50-90% fatty acid ester, triglyceride, polyol alkanoate, or dibasic fatty acid ester and 10-50% surfactant (monoalkyl ether of ethylene oxide-propylene oxide polymer). Thus, polyethylene-

having good thermal stability and low fume emission contain 50-90% fatty acid ester, triglyceride, polyol alkanoate, or dibasic fatty acid ester and 10-50% surfactant (monoalkyl ether of ethylene oxide-propylene oxide polymer). Thus, polyethylene-polypropylene glycol mono-2-ethylhexyl ether [64366-70-7] [smoke point 200°, volatility 7.2%, thin-film residue on stainless steel 1.6% (removable with hot soapy water)] was mixed in ratio 20:80, 30:70, and 40:60 with coconut oil to give spin finishes for polyester fibers. Aqueous emulsions (10 and 20%) of the finishes were stable for  $\geq$ 24 h.

IC D06M013-18; D06M013-16; C10M003-20

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finish fiber; surfactant finish fiber; lubricant
     finish fiber; coconut oil finish fiber;
     polyester fiber spin finish
IT
     Lubricants
        (esters, in spin finishes for fibers
IT
     Coconut oil
     RL: USES (Uses)
        (lubricant, in spin finishes for
IT
     Surfactants
        (polyoxyalkylene ethers, in spin finishes for
        fibers)
IT
     Polyester fibers, uses and miscellaneous
     Synthetic fibers
     RL: USES (Uses)
        (spin finishes for, lubricant-surfactant
        mixts. as)
ΙT
     Polyoxyalkylenes
     RL: USES (Uses)
        (spin finishes, containing lubricants, for
        polyester fibers)
IT
     9038-95-3
                 37311-00-5 64366-70-7
     RL: USES (Uses)
        (spin finishes, containing lubricants, for
        polyester fibers)
IT
     126-57-8
                31556-45-3
     RL: USES (Uses)
        (spin finishes, containing surfactants, for
        polyester fibers)
L133 ANSWER 53 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1981:67233
           Document No. 94:67233 A spin finish for
     synthetic fibers and methods of lubricating synthetic
     yarns with an aqueous emulsion containing this
     spin finish. Decker, Quintin William; Marcus,
     Erich; Koenig, Harvey Steven (Union Carbide Corp., USA). Eur. Pat.
     Appl. EP 17197 19801015, 81 pp. (English). CODEN: EPXXDW.
     APPLICATION: EP 1980-101687 19800328.
AB
     Polyethylene-polypropylene glycol alkylphenyl ether
     surfactants display good thermal stability, low fuming
     characteristics, and emulsification versatility when used to
     formulate spin finishes for synthetic
     fibers. Thus, treatment of nonylphenol with ethylene oxide
     (I) and propylene oxide (II) gave a colorless polyethylene-
     polypropylene glycol nonylphenyl ether [9082-01-3] (8 mol I
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CC

ST ·

40-7 (Textiles)

spin finish fiber; polyoxyalkylene

and 6.5 mol II) having smoke point 201°, volatility 2.4%/h, viscosity 322 and 150 cSt at 25°C and 100°F, resp., sp. gr. 1.023 at 25°, m.p. 7°, and cloud point 22°. The product had excellent heat stability and emulsification properties when tested with coconut oil, trimethylolpropane trispelargonate [126-57-8], and tridecyl stearate [31556-45-3]. C10M003-20; C10M003-22; D06M013-18; D06M013-20 39-8 (Textiles) surfactant emulsifier finish synthetic fiber; lubricant emulsifier finish synthetic fiber; polyethylene polypropylene glycol nonylphenyl ether Coconut oil RL: USES (Uses) (lubricants, spin finishes for synthetic fibers containing, low-fuming heat-stable polyethylenepolypropylene glycol alkylphenyl ethers as low-fuming heat-stable emulsifiers for) Emulsifying agents Surfactants (polyethylene-polypropylene glycol alkylphenyl ethers, heat-stable low-fuming, spin finishes containing lubricants and, for synthetic fibers) Lubricants (spin finishes for synthetic fibers containing, polyethylene-polypropylene glycol alkylphenyl ethers as low-fuming heat-stable emulsifiers for) Synthetic fibers RL: USES (Uses) (spin finishes for, containing lubricants and heat-stable low-fuming polyethylene-polypropylene glycol alkylphenyl ether emulsifiers) 9082-01-3 76483-16-4 RL: USES (Uses) (emulsifiers, heat-stable low-fuming, spin finishes containing lubricants and, for synthetic fibers) 126-57-8 31556-45-3 RL: USES (Uses) (lubricants, spin finishes for synthetic fibers containing, low-fuming heat-stable polyethylene-

IC CC

ST

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L133 ANSWER 54 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1977:586014 Document No. 87:186014 Surface coating of polymers.
Thompson, David Thomas; Jackson, Reginald; Ballard, Denis George Harold; Emmerson, James Derek; Morris, Stephen; Margetts, Anthony

polypropylene glycol alkylphenyl ethers as low-fuming

heat-stable emulsifiers for)

James; Gaitskell, John Nigel (Imperial Chemical Industries Ltd., UK). Brit. GB 1473667 19770518, 9 pp. (English). CODEN: BRXXAA. APPLICATION: GB 1974-10523 19750304.

Nylon 66 and poly(ethylene terephthalate) yarns were coated batchwise or continuously with polymers by treating the yarns with a Cr or Zr hydrocarbyl and exposing the treated yarns to monomer to cause it to polymerize onto the yarns. E.g., 3 g of polyester yarn without spin finish was dried overnight at 120°, cooled, immersed in 70 mL PhMe, treated with an excess 0.22mM tetrakis( $\pi$ -allyl)zirconium [12090-34-5] in PhMe, and exposed 2 h to 10 mL CH2:CHCN. A coating of polyacrylonitrile [25014-41-9] had formed on the yarn. The coated yarn was sandwiched between 2 150- $\mu$ -diameter stainless steel wires and the assembly embedded in polyethylene [9002-88-4]. The adhesive shear strength of the bond was 4.8 MN/m2 compared with 2.5 MN/m2 for a similar but uncoated yarn.

IT 9002-88-4

RL: USES (Uses)

(coating of, on synthetic **fibers**, catalysts for in situ polymerization in)

RN 9002-88-4 HCAPLUS

CN Ethene, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 74-85-1 CMF C2 H4

 $H_2C = CH_2$ 

IC C08F002-02

CC 39-8 (Textiles)

Section cross-reference(s): 42

coating process synthetic fiber; polyamide fiber coating polyethylene; polyester fiber coating polyolefin; polyethylene coating synthetic fiber; polyisoprene coating polyester fiber; polyacrylonitrile coating polyester fiber; catalyst polymer coating fiber; zirconium catalyst coating fiber; chromium catalyst coating fiber

IT Polymerization catalysts

(chromium and zirconium hydrocarbyls, for olefins in coating on polyamide and polyester fibers)

IT Polyamide fibers, uses and miscellaneous

RL: USES (Uses)

(coating of, with polyethylene, catalysts for in situ polymerization in)

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IT
     Polyester fibers, uses and miscellaneous
     RL: USES (Uses)
        (coating of, with polyolefins, catalysts for in situ
        polymerization in)
IT
     Coating process
        (of polyamide and polyester fibers, with
        polyolefins, catalysts for in situ polymerization in)
I \cdot T
     12082-46-1
     RL: CAT (Catalyst use); USES (Uses)
        (catalysts, for polymerization of acrylonitrile on polyester
        fiber)
IT
     12090-34-5
                  24356-01-2
     RL: CAT (Catalyst use); USES (Uses)
        (catalysts, for polymerization of olefins on
        synthetic fibers)
                 25014-41-9
IT
     9003-31-0
     RL: USES (Uses)
        (coating of, on polyester fiber, catalysts for in situ
       polymerization in)
IT
     9002-88-4
     RL: USES (Uses)
        (coating of, on synthetic fibers, catalysts for in situ
       polymerization in)
L133 ANSWER 55 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
           Document No. 84:32549 Polyamide yarn. Marshall, Robert
    Moore; Pak, Sung Kun; Dardoufas, Kimon C. (Allied Chemical Corp.,
    USA). U.S. US 3917893 19751104, 5 pp. (English). CODEN: USXXAM.
    APPLICATION: US 1974-438970 19740204.
AB
     Polyamide yarns were given a finish making them suitable for use in
    braided hose, rope, or drive belts, by treating with an aq
     . emulsion of an oxidized polyethylene [9002-88-4
     ] wax having average mol. weight 1000-3000 and containing 3-9% O,
    heating to 100-175°, and treating with a composition containing an
     aqueous dispersion of colloidal silica
     [7631-86-9] and a polyethylene glycol ester of a C6-18 aliphatic
    acid. Nylon 6 yarn(1260 denier/204 fibers) was treated
    with a water-resistant spin finish prepared by
    mixing 4 parts water with a composition consisting of 48.2% white mineral
    oil, 38.4% phosphated oleyl alc. ethoxylated with .apprx.7 mols of
    ethylene oxide, 6.6% sulfonated succinic acid ester of oleyl alc.,
     and 6.8% oxidized polyethylene wax. The yarn was overfinished with
    a composition containing 30 parts of polyethylene glycol monopelargonate
     [31621-91-7] and 70 parts of a commercially available
    colloidal aqueous silica dispersion containing
    30 weight% solids to give a yarn with 0.3% silica and could
    be processed through a coning operation and was suitable for
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braiding and winding operations during manufacture of industrial

rope. IC D02J; D01H

INCL 428395000

CC 39-10 (Textiles)

IT Polyamide fibers RL: USES (Uses)

(finishing agents for yarns of, for improved coning and braiding)

IT Ethene, homopolymer, oxidized

RL: USES (Uses)

(wax, polyamide yarn finished by, for improved coning, and braiding)

L133 ANSWER 56 OF 56 HCAPLUS COPYRIGHT 2005 ACS on STN
1970:56306 Document No. 72:56306 Polyolefin foams. Hosoda,
Kirokuro; Nakae, Hiroyuki; Shiina, Naonori; Kadowaki, Yoshio
(Furukawa Electric Co., Ltd.). Ger. Offen. DE 1930134 19691218, 43
pp. (German). CODEN: GWXXBX. APPLICATION: DE 1969-1930134
19690613.

Polyolefin foams containing 70-99.5 weight% polyolefin AB and 0.5-30 weight % polybutadiene (I) are prepared by heating a mixture of the 2 polymers in the presence of a blowing agent. No crosslinking agent is needed. The composition can be foamed in a continuous process by extrusion, injection molding, or blow molding. Light materials with high flexural strength, particularly useful as building materials, are obtained by addition of I to the polyolefins. Thus, high-d. polyethylene (II) with a melt index of 0.5 and a d. of 0.96 g/cm3 was mixed 2 min with I containing ≥98% cis-1,4 bonds on a roll mill at 140-5°. To this moxt. was added 5 parts 1,1'-azobisformamide, mixing was continued 15 min, the mixture was hot pressed at 140-5° into a plate 7 mm thick, cut into a 40 + 40-mm piece, the piece was wrapped in Al foil, and foamed 10 min at 200° in a bath of Wood's alloy. The product had the following properties (parts I, parts II, d., thickness of the foamed plate in mm, compression elasticity for 25% compression in kg/cm2 given): 0,100, 0.43, 8, -; 1, 99, 0.20, 19, -; 5, 95, 0.10, 30, -; 10, 90, 0.11, 31, 4.3; 20, 80, 0.11, -, 3.6; 30, 70, 0.12, -, 3.0; 40, 60, 0.15, -, 1.4; 70, 30, 0.45, -, -. Other olefins used were low-d. II, isotactic polypropylene, an ethylene-vinyl acetate copolymer, and chlorosulfonated polyethylene; an ethylene-acrylic acid copolymer, an ethylene-propylene copolymer , an ethylene-butene copolymer, chlorinated polyethylene or polypropylene, and poly(4-methyl-1-pentene) are also The foamable mixture may also contain Zn stearate, Al stearate, Ca silicate, CaCO3, ZnO, talc, TiO2, or diatomaceous earth as foam nucleating agent and a thickening agent such as glass fibers or carbon black. .

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IT
     9002-88-4, uses and miscellaneous
     RL: USES (Uses)
        (cellular, containing butadiene polymers)
     9002-88-4 HCAPLUS
RN
CN
     Ethene, homopolymer (9CI) (CA INDEX NAME)
     CM
          1
     CRN 74-85-1
     CMF C2 H4
H_2C = CH_2
     RL: USES (Uses)
        (chlorosulfonated, cellular, contg. butadiene polymers
IC
CC
     36 (Plastics Manufacture and Processing)
ST
     polyolefin polybutadiene blend foams; polybutadiene
     polyolefin blend foams; foams polyolefins
     polybutadiene blend; polyethylene polybutadiene blend foams;
     polypropylene polybutadiene blend foams
IT
     Olefins, uses and miscellaneous
     RL: USES (Uses)
        (cellular, containing butadiene polymers)
IT
     9003-17-2, uses and miscellaneous
     RL: USES (Uses)
        (cellular olefin polymers containing)
IT
     9002-88-4, uses and miscellaneous 9003-07-0, uses and
     miscellaneous 24937-78-8, uses and miscellaneous
     RL: USES (Uses)
        (cellular, containing butadiene polymers)
IT
     9002-88-4, uses and miscellaneous
     RL: USES (Uses)
        (chlorosulfonated, cellular, containing butadiene polymers)
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